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(54) Title: SMALL STREPTOCOCCUS PYOGENES ANTIGENS AND THEIR USE

(57) Abstract: The present invention relates to a peptide consisting of one antigen of Streptococcus pyogenes (S. pyogenes) of any of the SEQ ID NOS: 1 to 7 or a functional active variant thereof, optionally further consisting of additional amino acid residue(s); a nucleic acid coding for the same; a pharmaceutical composition, especially a vaccine, comprising said peptide or said nucleic acid; an antibody or functional active fragment thereof specifically binding to the antigen; a hybridoma cell line which produces said antibody; a method for producing said antibody; a pharmaceutical composition comprising said antibody; the use of said peptide or said nucleic acid for the manufacture of a medicament for the immunization or treatment of a subject; the use of said antibody or functional fragment thereof for the manufacture of a medicament for the treatment of an infection; a method of diagnosing a S. pyogenes infection; a method for identifying a ligand capable of binding to said peptide; and the use of said peptide for the isolation and/or purification and/or identification of an interaction partner of the peptide.



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Small Streptococcus pyogenes Antigens and their Use

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The present invention relates to a peptide consisting of one antigen of *Streptococcus pyogenes* (S. pyogenes) of any of the SEQ ID NOS: 1 to 7 or a functional active variant thereof, optionally further consisting of additional amino acid residue(s); a nucleic acid coding for the same; a pharmaceutical composition, especially a vaccine, comprising said peptide or said nucleic acid; an antibody or functional active fragment thereof specifically binding to the antigen; a hybridoma cell line which produces said antibody; a method for producing said antibody; a pharmaceutical composition comprising said antibody; the use of said peptide or said nucleic acid for the manufacture of a medicament for the immunization or treatment of a subject; the use of said antibody or functional fragment thereof for the manufacture of a medicament for the treatment of an infection; a method of diagnosing a S. pyogenes infection; a method for identifying a ligand capable of binding to said peptide; and the use of said peptide for the isolation and/or purification and/or identification of an interaction partner of the peptide.

Streptococcus pyogenes, also called group A streptococcus (GAS), is an important grampositive extracellular bacterial pathogen and commonly infects humans. GAS colonizes the throat or skin and is responsible for a number of suppurative infections and non-suppurative sequelae. It is primarily a disease of children and causes a variety of infections including bacterial pharyngitis, scarlet fever, impetigo and sepsis in humans. Decades of epidemiological studies have led to the concept of distinct throat and skin strains, where certain serotypes are often associated with throat or skin infections, respectively (Cunningham, M. (2000). Clin Microbiol Rev 13: 470-511). GAS has been discovered responsible for streptococcal toxic shock syndrome associated necrotizing fasciitis which is recently resurgent in the USA (Cone, L., et al. (1987). New Engl J Med 317: 146-9; Stevens, D. (1992). Clin Infect Dis 14: 2-11) and has been described as the "flesh eating" bacterium which invades skin and soft tissues leading to tissue or limb destruction.

Several post-streptococcal sequelae may occur in humans subsequent to infection, such as acute rheumatic fever, acute glomerulonephritis and reactive arthritis. Acute rheumatic

fever and rheumatic heart disease are of these the most serious autoimmune sequelae and have led to disability and death of children worldwide. *S. pyogenes* can also causes severe acute diseases such as scarlet fever and necrotizing fasciitis and has been associated with Tourette's syndrome, tics and movement and attention disorders.

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Group A streptococci are the most common bacterial cause of sore throat and pharyngitis and account for at least 16% of all office calls in a general medical practice, season dependent (Hope-Simpson, R. (1981). J Hyg (Lond) 87: 109-29). It primarily affects children in school-age between 5 to 15 years of age (Cunningham, supra). All ages are susceptible to spread of the organism under crowded conditions, for example in schools. GAS are not considered normal flora though, but pharyngeal carriage of group A streptococci can occur without clinical symptoms.

Group A streptococci can be distinguished by the Lancefield classification scheme of serologic typing based on their carbohydrate or classified into M protein serotypes based on a surface protein that can be extracted by boiling bacteria with hydrochloric acid. This has led to the identification of more than 80 serotypes, which can also be typed by a molecular approach (emm genes). Molecular typing has identified more than 150 individual emm types. Certain M protein serotypes of *S. pyogenes* are mainly associated with pharyngitis and rheumatic fever, while others mainly seem to cause pyoderma and acute glomerulonephritis (Cunningham, supra).

Also implicated in causing pharyngitis and occasionally toxic shock are group C and G streptococci, which must be distinguished after throat culture (Hope-Simpson, supra; Bisno, A., et al. (1987). <u>Infect Immun</u> 55: 753-7).

Currently, streptococcal infections can only be treated by antibiotic therapy. However, 25-30% of those treated with antibiotics show recurrent disease and/or shed the organism in mucosal secretions. There is at present no preventive treatment (vaccine) available to avoid streptococcal infections.

Thus, there remains a need for an effective treatment to prevent or ameliorate streptococcal infections. A vaccine could not only prevent infections by streptococci, but more

specifically prevent or ameliorate colonization of host tissues, thereby reducing the incidence of pharyngitis and other suppurative infections. Elimination of non-suppurative sequelae such as rheumatic fever, acute glomerulonephritis, sepsis, toxic shock and necrotizing fasciitis would be a direct consequence of reducing the incidence of acute infection and carriage of the organism. Vaccines capable of showing cross-protection against other streptococci would also be useful to prevent or ameliorate infections caused by all other beta-hemolytic streptococcal species, namely groups A, B, C and G.

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A vaccine can contain a whole variety of different antigens. Examples of antigens are whole-killed or attenuated organisms, subfractions of these organisms/tissues, proteins, or, in their most simple form, peptides. Antigens can also be recognized by the immune system in form of glycosylated proteins or peptides and may also be or contain polysaccharides or lipids. Short peptides can be used since for example cytotoxic T-cells (CTL) recognize antigens in form of short usually 8-11 amino acids long peptides in conjunction with major histocompatibility complex (MHC). B-cells can recognize linear epitopes as short as 4-5 amino acids, as well as three-dimensional structures (conformational epitopes).

In some circumstances, adjuvants may be useful for sustaining antigen-specific immune responses. Primarily, adjuvants are acting, but are not restricted in their mode of action, on so-called antigen presenting cells (APCs). These cells usually first encounter the antigen(s) followed by presentation of processed or unmodified antigen to immune effector cells. Intermediate cell types may also be involved. Only effector cells with the appropriate specificity are activated in a productive immune response. The adjuvant may also locally retain antigens and co-injected other factors. In addition the adjuvant may act as a chemoattractant for other immune cells or may act locally and/or systemically as a stimulating agent for the immune system.

Approaches to develop a group A streptococcal vaccine have focused mainly on the cell surface M protein of S. pyogenes (Bessen, D., et al. (1988). <u>Infect Immun</u> 56: 2666-2672; Bronze, M., et al. (1988). <u>J Immunol</u> 141: 2767-2770). Since more than 80 different M serotypes of S. pyogenes exist and new serotypes continually arise (Fischetti, V. (1989). Clin Microbiol Rev 2: 285-314), inoculation with a limited number of serotype-specific M

protein or M protein derived peptides will not likely be effective in protecting against all other M serotypes. Furthermore, it has been shown that the conserved region of the M protein contains an amino acid sequence, which is immunologically cross-reactive with human heart tissue, which is thought to account for heart valve damage associated with rheumatic fever (Fenderson, P., et al. (1989). <u>J Immunol</u> 142: 2475-2481).

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There are other proteins under consideration for vaccine development, such as the erythrogenic toxins, streptococcal pyrogenic exotoxin A and streptococcal pyrogenic exotoxin B (Lee, P. K. (1989). <u>J Clin Microbiol</u> 27: 1890-2). Immunity to these toxins could possibly prevent the deadly symptoms of streptococcal toxic shock, but it may not prevent colonization by group A streptococci.

The use of the above described proteins as antigens for a potential vaccine as well as a number of additional candidates (Ji, Y., et al. (1997). <u>Infect Immun</u> 65: 2080-2087; Guzman, C., et al. (1999). <u>J Infect Dis</u> 179: 901-6) resulted mainly from a selection based on easiness of identification or chance of availability. There is a demand to identify efficient and relevant antigens for *S. pyogenes*.

WO 2004/078907 describes a method for identification, isolation and production of hyperimmune serum reactive antigens from *Streptococcus pyogenes*.

The antigens described herein focus on regions shown in the present application to be protective. A suitable antigen size to obtain protection varies based on different factors such as the type of protective epitope (e.g., conformational versus linear) and the number of protective epitopes providing a level of protection. Large antigens containing regions not providing useful protection may be disadvantageous in the context of immunization. First, providing of smaller antigens eases production of the protein in recombinant form. It is generally accepted that it is more difficult to produce larger proteins. Smaller proteins may be produced in a more economic manner, thus saving costs, particularly in the health care system. Second, reducing the size of antigenic proteins used for vaccination may lead to safer products. Eliminating extra sequences in antigenic proteins is desirable, since this reduces the probability of inducing antibodies which can cause cross-reactions with human tissues. Third, proteins used for vaccination may contain more than one antigen, the

antigens directed either against the same disease or against different diseases, in order to

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obtain a more effective vaccination or vaccination against several diseases. However, if the single antigens are too large a combination into one protein is not feasible.

Accordingly, one problem underlying the present invention was to provide alternative 5 means for the development of medicaments such as vaccines against S. pyogenes infection, particularly smaller proteins.

Surprisingly, the object has been solved by a peptide consisting of one antigen of S. pyogenes of the SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7 or a functional active variant of one antigen of S. pyogenes of the SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7. These peptides are referred to as antigenic peptides.

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The sequences of SEQ ID NOS: 1 to 7 are characterized in table 1 of the present specification. The underlying amino acid sequences are disclosed in the attached sequence data. The peptides of SEQ ID NOS: 1 to 7 have been shown to induce an immune response and/or to show protection against S. pyogenes in a sepsis and/or lethality model (see Example 1). Functional active variants are obtained by changing the sequence of the antigen as defined below and are characterized by having a biological activity similar to that displayed by the antigen of any of the sequences of SEQ ID NO: 1 to 7 from which it is derived, including the ability to induce immune responses and/or to show protection against S. pyogenes e.g. in a sepsis and/or lethality model.

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In some embodiments of the invention the peptide of the invention consists of one antigen of S. pyogenes of the SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7 or a functional active variant of one antigen of S. pyogenes of the SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7; and

1 to 350 additional amino acid residue(s), preferably 1 to 200, more preferably 1 to a) 150, even more preferably at most 1 to 100, still more preferably at most 1 to 50,

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the antigen is SEQ ID NO: 1; or

1 to 200 additional amino acid residue(s), preferably 1 to 150, more preferably 1 to b) 100, even more preferably at most 1 to 50, still more preferably at most 1 to 25, most preferably 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 additional amino acids residue(s) if the antigen is SEQ ID NO: 2; or

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- 1 to 100 additional amino acid residue(s), preferably 1 to 75, more preferably 1 to c) 50, even more preferably at most 1 to 25, still more preferably at most 1 to 10, most preferably 1, 2, 3, 4 or 5 additional amino acids residue(s) if the antigen is that of SEQ ID NO: 3; or
- 1 to 150 additional amino acid residue(s), preferably 1 to 100, more preferably 1 to d) 75, even more preferably at most 1 to 50, still more preferably at most 1 to 25, most preferably 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 additional amino acids residue(s) if the antigen is that of SEQ ID NO: 4; or
- 1 to 450 additional amino acid residue(s), preferably 1 to 300, more preferably 1 to 15 e) 150, even more preferably at most 1 to 100, still more preferably at most 1 to 50, most preferably 1, 2, 3, 4, 5, 10, 20, 30 or 40 additional amino acids residue(s) if the antigen is SEQ ID NO: 5; or
- 1 to 250 additional amino acid residue(s), preferably 1 to 200, more preferably 1 to **f**) 150, even more preferably at most 1 to 100, still more preferably at most 1 to 50, 20 most preferably 1, 2, 3, 4, 5, 10, 15, 20 or 25 additional amino acids residue(s) if the antigen is SEQ ID NO: 6 or SEQ ID NO: 7.

The antigen of S. pyogenes can be any of the antigens as defined above, namely as defined in any of the SEQ ID NOS: 1, 2, 3, 4, 5, 6 or 7, or a functional active variant thereof, 25 wherein the functional active variant is as defined below.

The antigen or the functional active variant thereof may have added at least one additional amino acid residue heterologous or homologous to the peptide. Homologous refers to any amino acid or amino acid sequence which is identical to the amino acid sequence of the S. pyogenes protein from which the antigen is derived, wherein the sequences of SEQ ID NO: 1 to 7 are derived from the following proteins:

Sequence	derived from protein (as disclosed in e.g. WO 2004/078907 or in the attached sequence data)	
SEQ ID NO: 1	Spy0269	
SEQ ID NO: 2	Spy0292	
SEQ ID NO: 3	Spy0292	
SEQ ID NO: 4	Spy0416	
SEQ ID NO: 5	Spy0416	
SEQ ID NO: 6	Spy0416	
SEQ ID NO: 7	Spy0872	

In one embodiment the antigen or the functional active variant thereof having one or more additional amino acid residues (see above, particularly as defined in items (a) to (f)) further encompasses at least one amino acid residue heterologous to the antigen. The feature "heterologous amino acid" or "amino acid heterologous to the antigen or protein" refers to any amino acid which is different from that amino acid located adjacent to the antigen or protein in any naturally occurring protein of *S. pyogenes*, particularly from that of *S. pyogenes* SF370 (serotype M1). Therefore, the protein of the invention encompassing at least one heterologous amino acid refers to a protein which is different from any naturally occurring protein of *S. pyogenes* or fragment thereof, particularly which is different from that of *S. pyogenes* SF370 (serotype M1). The proteins from which the antigens of the invention are derived as well as a reference for their sequences are listed above.

In certain embodiments, the peptide consists of the antigen, optionally the at least one additional amino acid residue as defined above, and at least one additional heterologous amino acid sequence comprising a marker protein.

The additional sequence or amino acid residue(s) as defined above consists of (an) amino acid residue(s), which may be any amino acid, which may be either an L-and/or a D-amino acid, naturally occurring and otherwise. Preferably the amino acid is any naturally occurring amino acid such as alanine, cysteine, aspartic acid, glutamic acid, phenylalanine, glycine, histidine, isoleucine, lysine, leucine, methionine, asparagine, proline, glutamine, arginine, serine, threonine, valine, tryptophan or tyrosine.

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However, the amino acid residue(s) may also be (a) modified or unusual amino acid(s). Examples of those are 2-aminoadipic acid, 3-aminoadipic acid, beta-alanine, 2-aminobutyric acid, 4-aminobutyric acid, 6-aminocaproic acid, 2-aminoheptanoic acid, 2-aminoisobutyric acid, 3-aminoisobutyric acid, 2-aminopimelic acid, 2,4-diaminobutyric acid, desmosine, 2,2'-diaminopimelic acid, 2,3-diaminopropionic acid, N-ethylglycine, N-ethylasparagine, hydroxylysine, allo-hydroxylysine, 3-hydroxyproloine, 4-hydroxyproloine, isodesmosine, allo-isoleucine, N-methylglycine, N-methylisoleucine, 6-N-Methyllysine, N-methylvaline, norvaline, norleucine or ornithine.

Additionally, the amino acid(s) may be subject to modifications such as posttranslational modifications. Examples of modifications include acetylation, amidation, blocking, formylation, γ-carboxyglutamic acid hydroxylation, glycosilation, methylation, phosphorylation and sulfatation.

If more than one additional or heterologous amino acid residue is present in the peptide, the amino acid residues may be the same or different from one another.

The antigenic peptide may be flanked by the amino acid residue(s) C-terminally, N-terminally, or C- and N-terminally.

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In a further embodiment the peptide is as described above in the different embodiments, and contains a region that is essentially identical to any of the antigens of the SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7, but differs from the antigens of any of the of the SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7, in that is it derived from a homologous sequence of a different serotype of *S. pyogenes*, particularly wherein the serotype is M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370.

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Accordingly, the present invention also relates to antigens of different *S. pyogenes* isolates. Such homologues may easily be identified and isolated based on the nucleic acid and amino acid sequences disclosed herein. A homologous antigen of a different serotype may

be identified by e.g. sequence alignment. The homologous antigen sequence may vary from the antigen of any of the sequences of SEQ ID NO: 1 to 7 by one or more amino acid substitutions, deletions and/or additions. Preferably the homologous antigen sequence has the sequence of any of the homologous variants identified in the attached listing of amino acid sequences.

Examples of homologous sequences of a different serotype are detailed in the attached sequence data. Particularly, sequences homologous to the respective peptide of the invention are those listed below:

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Full length amino acid sequence (SEQ ID NO)	Peptide of the invention (SEQ ID NO)	Homologous amino acid sequen- ces (SEQ ID NOS)
57	1	58 to 67
68	2	69 to 78
68	3	79 to 88
89	4	90 to 99
89	5	100 to 109
89	6	110 to 119
120	7	121 to 130

There are more than 150 emm types distinguished to date and the typing is based on the variable region at the 5' end of the emm gene (see e.g. Vitali, L., et al. (2002) <u>J. Clin. Microbiol</u> 40: 679-681). The presence of a homologous antigen can accordingly be determined for every emm type. In addition it is possible to determine the variability of a particular antigen in the various emm types as described for the *sic* gene (Hoe N., et al. (2001) <u>J. Inf. Dis.</u> 183: 633-9). The influence of the various M serotypes on the kind of disease it causes is summarized in a recent review (Cunningham, supra). In particular, two groups of serotypes can be distinguished:

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- 1) Those causing Pharyngitis and Scarlet fever (e.g. M types 1, 3, 5, 6, 14, 18, 19, 24)
- 2) Those causing Pyoderma and Streptococcal skin infections (e.g. M types 2, 49, 57, 59, 60, 61)

This can serve as the basis to identify the relevance of an antigen for the use as a vaccine or in general as a drug targeting a specific disease.

The information e.g. from the homepage of the Centers for Disease Control and Prevention (CDC) (http://www.cdc.gov/ncidod/biotech/strep/emmtypes.htm) gives a dendrogram showing the relatedness of various emm types. Further relevant references are Vitali et al., supra (molecular emm typing method), Enright et al., Infection and Immunity 69: 2416-2427. (2001) (alternative molecular typing method (MLST)), Hoe et al., supra (example for the variation of one antigen (*sic*) in many different serotypes) and Cunningham, supra (review on GAS pathogenesis). All emm types are completely listed and are available at publicly available databases (e.g., through the CDC).

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In another embodiment of the present invention the variant is a fragment. The fragment is characterized by being derived from the antigen as defined above by one or more amino acid deletions. The deletion(s) may be C-terminally, N-terminally and/or internally. Preferably the fragment is obtained by at most 10, 20, 30, 40, 50, 60, 80, 100, 150 or 200, more preferably by at most 10, 20, 30, 40 or 50, even more preferably at most 5, 10 or 15, still more preferably at most 5 or 10, most preferably 1, 2, 3, 4 or 5 amino acid deletion(s). The functional active fragment of the invention is characterized by having a biological activity similar to that displayed by the complete antigen, including the ability to induce immunization and/or to show protection against S. pyogenes e.g. in a sepsis and/or lethality model. The fragment of an antigen is functional active in the context of the present invention, if the activity of the fragment amounts to at least 10%, preferably at least 25%, more preferably at least 50%, even more preferably at least 70%, still more preferably at least 80%, especially at least 90%, particularly at least 95%, most preferably at least 99% of the activity of the antigen without sequence alteration. These fragments may be designed or obtained in any desired length, including as small as about 50 to 80 amino acids in length.

The functional active fragment may be also characterized by other structural features. Accordingly, in one preferred embodiment of the invention the functional active fragments consists of at least 60%, preferably at least 70%, more preferably at least 80%, still more

preferably at least 90%, even more preferably at least 95%, most preferably 99% of the amino acids of the antigen of any of the SEQ ID NOS: 1 to 7. The functional active fragment as defined above may be derived from the peptide by one or more amino acid deletions. The deletions may be C-terminally, N-terminally and/or internally.

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Another preferred embodiment of the invention relates to a peptide as defined above in the previous embodiments, wherein the antigen is a functional active variant of an antigen of any of the SEQ ID NOS: 1 to 7 and wherein the variant has at least 50% sequence identity to the antigen of any of the SEQ ID NOS: 1 to 7. In a more preferred embodiment the functional active variant has a sequence identity of at least 60%, preferably at least 70%, more preferably at least 80%, still more preferably at least 90%, even more preferably at least 95%, most preferably 99% to the antigen of any of the SEQ ID NOS: 1 to 7.

The percentage of sequence identity can be determined e.g. by sequence alignment. Methods of alignment of sequences for comparison are well known in the art. Various programs and alignment algorithms have been described e.g. in Smith and Waterman, Adv. Appl. Math. 2: 482, 1981 or Pearson and Lipman, Proc. Natl. Acad. Sci. U.S.A. 85: 2444-2448, 1988.

The NCBI Basic Local Alignment Search Tool (BLAST) (Altschul et al., J. Mol. Biol. 215: 403-410, 1990) is available from several sources, including the National Center for Biotechnology Information (NCBI, Bethesda, MD) and on the Internet, for use in connection with the sequence analysis programs blastp, blastn, blastx, tblastn and tblastx. Variants of an antigen of any of the sequences of SEQ ID NOS: 1 to 7 are typically characterized using the NCBI Blast 2.0, gapped blastp set to default parameters. For comparisons of amino acid sequences of at least 35 amino acids, the Blast 2 sequences function is employed using the default BLOSUM62 matrix set to default parameters, (gap existence cost of 11, and a per residue gap cost of 1). When aligning short peptides (fewer than around 35 amino acids), the alignment is performed using the Blast 2 sequences function, employing the PAM30 matrix set to default parameters (open gap 9, extension gap 1 penalties). Methods for determining sequence identity over such short windows such as 15 amino acids or less are described at the website that is maintained by the National

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Center for Biotechnology Information in Bethesda, Maryland (http://www.ncbi.nlm.nih.gov/BLAST/).

The functional active variant of an antigen is obtained by sequence alterations in the antigen, wherein the antigen with the sequence alterations retains a function of the unaltered antigen, e.g. having a biological activity similar to that displayed by the complete antigen, including the ability to induce an immune response and/or to show protection against *S. pyogenes* e.g. in a sepsis and/or lethality model. Such sequence alterations can include, but are not limited to, conservative substitutions, deletions, mutations and insertions. These characteristics of the functional active variant can be assessed e.g. as detailed in Example 1. In the context of the present invention a variant specifically has a biological activity similar to that displayed by the antigen without alteration, including the ability to induce an immune response and/or to show protection against *S. pyogenes* e.g. in a sepsis and/or lethality model if the activity of the variant amounts to at least 10%, preferably at least 25%, more preferably at least 50%, even more preferably at least 70%, still more preferably at least 80%, especially at least 90%, particularly at least 95%, most preferably at least 99% of the activity of the antigen without sequence alterations.

The term "functional active variant" includes naturally-occurring allelic variants, as well as mutants or any other non-naturally occurring variants. As is known in the art, an allelic variant is an alternate form of a (poly)peptide that is characterized as having a substitution, deletion, or addition of one or more amino acids that does essentially not alter the biological function of the polypeptide. By "biological function" is meant a function of the polypeptide in the cells in which it naturally occurs, even if the function is not necessary for the growth or survival of the cells. For example, the biological function of a porin is to allow the entry into cells of compounds present in the extracellular medium. The biological function is distinct from the antigenic function. A polypeptide can have more than one biological function.

Within any species of the living world, allelic variation is the rule. For example, any bacterial species, e.g. S. pyogenes, is usually represented by a variety of strains (characterized by clonal reproduction) that differ from each other by minor allelic variations. Indeed, a polypeptide that fulfils the same biological function in different

strains can have an amino acid sequence that is not identical in each of the strains. Such an allelic variation is equally reflected at the polynucleotide level.

Allelic variation is very common within the *S. pyogenes* species. Such allelic variation is also the basis for the molecular typing of group A streptococcal strains by emm typing as described above (see, e.g. Facklam, R. et al. (1999) Emerg Infect Dis. 5: 247-53 or http://www.cdc.gov/ncidod/biotech/strep/emmtypes.htm). Further, genes such as *sic* are subject to allelic variation (Hoe N., et al. (2001) J. Inf. Dis. 183: 633-9). However, proteins with large allelic variation are in general not suitable candidates for a vaccine, as immunization would not protect against infection with all strains, or alternative immunization would possibly induce the emergence of new allelic variants not covered by the vaccine.

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In a preferred embodiment, the functional active variant or fragment derived from the antigen by amino acid exchanges, deletions or insertions may also conserve, or more preferably improve, the activity (as defined above). Furthermore, these peptides may also cover epitopes, which trigger the same or preferably an improved T cell response. These epitope are referred to as "heteroclitic". They have a similar or preferably greater affinity to MHC/HLA molecules, and the ability to stimulate the T cell receptors (TCR) directed to the original epitope in a similar or preferably stronger manner. Heteroclitic epitopes can be obtained by rational design i. e. taking into account the contribution of individual residues to binding to MHC/HLA as for instance described by (Rammensee, H. et al., 1999, Immunogenetics. 50: 213-219), combined with a systematic exchange of residues potentially interacting with the TCR and testing the resulting sequences with T cells directed against the original epitope. Such a design is possible for a skilled man in the art without much experimentation.

In a still more preferred embodiment of the invention the functional active variant of an antigen of any of the SEQ ID NOS: 1 to 7 having at least 50% sequence identity to the antigen of any of the SEQ ID NOS: 1 to 7, especially at least 60%, preferably at least 70%, more preferably at least 80%, still more preferably at least 90%, even more preferably at least 95%, most preferably 99% to the antigen of any of the SEQ ID NOS: 1 to 7 is derived from the antigen of any of the sequences of SEQ ID NOS: 1 to 7 by conservative

substitutions. Conservative substitutions are those that take place within a family of amino acids that are related in their side chains and chemical properties. Examples of such families are amino acids with basic side chains, with acidic side chains, with non-polar aliphatic side chains, with non-polar aromatic side chains, with uncharged polar side chains, with small side chains, with large side chains etc.. In one embodiment, one conservative substitution is included in the peptide. In another embodiment, two conservative substitutions or less are included in the peptide. In a further embodiment, three conservative substitutions or less are included in the peptide.

Examples of conservative amino acid substitutions include, but are not limited to, those listed below:

	Original Residue	Conservative Substitutions
	Ala	Ser
15	Arg	Lys
	Asn	Gln; His
	Asp	Glu
	Cys	Ser
	Gln	Asn
20	Glu	Asp
	His	Asn; Gln
	Ile	Leu, Val
	Leu	Ile; Val
	Lys	Arg; Gln; Asn
25	Met	Leu; Ile
	Phe	Met; Leu; Tyr
	Ser	Thr
	Thr	Ser
	Trp	Tyr
30	Tyr	Trp; Phe
	Val	Ile; Leu

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Examples of suitable variants of the peptide of the invention obtained by one or more amino acid exchange(s), deletion(s) and/or insertion(s) may be derived from data provided in tables 5 to 7 and 9. Particularly, tables 5 to 7 and 9 list naturally occurring amino acid alterations (substitutions, insertions, deletions) at particular positions in comparison to *S. pyrogenes* SF370.

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With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 1, the variant of the invention may differ from the peptide having SEQ ID NO: 1 by one or more of the alterations identified in table 5.

With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 2, the variant of the invention may differ from the peptide having SEQ ID NO: 2 by one or more of the alterations identified in table 6.

With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 3, the variant of the invention may differ from the peptide having SEQ ID NO: 3 by one or more of the alterations identified in table 6.

With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 4, the variant of the invention may differ from the peptide having SEQ ID NO: 4 by one or more of the alterations identified in table 7.

With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 5, the variant of the invention may differ from the peptide having SEQ ID NO: 5 by one or more of the alterations identified in table 7.

With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 6, the variant of

the invention may differ from the peptide having SEQ ID NO: 6 by one or more of the alterations identified in table 7.

With respect to a variant of a peptide having (i.e. consisting of or comprising as defined above, particularly as defined in the above items (a) to (f)) SEQ ID NO: 7, the variant of the invention may differ from the peptide having SEQ ID NO: 7 by one or more of the alterations identified in table 9.

It should be understood that variants obtained from a peptide of the invention by one or more sequence alterations in accordance with tables 5 to 7 and 9 are preferred.

A further aspect of the present invention describes a peptide comprising an amino acid sequence with at least 95% sequence identity to at least one of SEQ ID NO: 1, 2, 3, 4, 5, 6 or 7. In different embodiment the peptide comprises, consists, or consists essentially of a region of at least 95%, at least 97% or at least 99% identical to SEQ ID NO: 1, 2, 3, 4, 5, 6 or 7, or differs by 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 amino acid alteration(s). In one embodiment the term "consist" may be as defined in the above items (a) to (f)). Preferably, the peptide does not contain a full-length naturally occurring Spy0269, Spy0292, Spy0416A (amino acids 33-867), or Spy0872.

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SEQ ID NO: 1, 2, 3, 4, 5, 6 or 7 provide core sequences useful for producing a protective immune response. SEQ ID NO: 1 provides an amino acid core from amino acids 37-488 of Spy0269. SEQ ID NO: 2 provides a core region of amino acids 23-184 of Spy0292. SEQ ID NO: 3 provides a core of amino acids 23-300 of Spy0292, which is a longer-length sequence containing the shorter-length core sequence of 23-184 of Spy0292 provided in SEQ ID NO: 2. Surprisingly, the shorter fragment Spy0292-1 (SEQ ID NO: 2) shows even greater protection in the mouse model compared to the longer fragment Spy0292-3 (SEQ ID NO: 3), as depicted in Figure 1. As described above, smaller peptides are in general advantageous over larger ones, since they may be produced in a more economic manner, they reduce the probability of inducing antibodies which can cause cross-reactions with human tissues, and they facilitate the preparation of combination vaccines comprising more than one antigen. SEQ ID NO: 4, 5, and 6 provide different Spy0416A core sequences of varying activity. SEQ ID NO: 5 provides a common core of amino acids 148-

458 of Spy0416A and has the lowest activity. SEQ ID NO: 6 provides a core sequence containing amino acids 72-558 of Spy0416A with greater activity than the shorter core. SEQ ID NO: 4 provides an amino acid core containing amino acids 34-677 of Spy0416, also with activity greater than the 148-458 core.

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Based on the guidance provided herein different peptides can be designed taking into account the core sequences provided in SEQ ID NOs: 1-7. Such guidance includes structurally related peptides containing (1) internal alterations; (2) additional amino acid groups at the amino and/or carboxyl terminus; and/or (3) additional modification(s) as described herein.

For structurally related peptides, each amino acid alteration is independently either an addition, substitution, or deletion. In a further embodiment, the amino terminus is methionine. The presence of methionine may be useful for recombinant expression. In some cases, the methionine may be initially present as a result of translation and subsequently cleaved. Additional examples and embodiments, including broader embodiments and some further descriptions applicable for structurally related peptides such as functional variants are provided above, particularly in the description of functional active variants.

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In another subject of the invention the peptide as described above comprises or consists of at least 2, preferably at least 3, more preferably at least 4 antigens as defined above. If two ore more peptides derived from the same full length sequence (e.g Spy0292 or Spy0416) are combined into one peptide, these sequences do preferably not overlap. In one embodiment the term "consist" may be as defined in the above items (a) to (f)).

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In another embodiment of the invention the peptide as defined above may be modified by one or more of a variety of chemical techniques to produce derivatives having essentially the same activity (as defined above for fragments and variants) as the modified peptides, and optionally having other desirable properties. For example, carboxylic acid groups of the protein, whether C-terminal or side chain, may be provided in the form of a salt of a pharmaceutically-acceptable cation or esterified to form an ester, or converted to an amide. Amino groups of the peptide, whether amino-terminal or side chain, may be in the form of

a pharmaceutically-acceptable acid addition salt, such as the HCl, HBr, acetic, benzoic, toluene sulfonic, maleic, tartaric and other organic salts, or may be converted to an amide. Hydroxyl groups of the peptide side chains may be converted to alkoxy or to an ester using well recognized techniques. Phenyl and phenolic rings of the peptide side chains may be substituted with one or more halogen atoms, such as fluorine, chlorine, bromine or iodine, or with alkyl, alkoxy, carboxylic acids and esters thereof, or amides of such carboxylic acids. Thiols can be protected with any one of a number of well recognized protecting groups, such as acetamide groups.

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Peptides of this invention may be in combination with outer surface proteins or other proteins or antigens of other proteins. In such combination, the antigen may be in the form of a fusion protein. The antigen of the invention may be optionally fused to a selected peptide or protein derived from other microorganisms. For example, an antigen or polypeptide of this invention may be fused at its N-terminus or C-terminus to a polypeptide from another pathogen or to more than one polypeptide in sequence. Peptides which may be useful for this purpose include polypeptides identified by the prior art.

In an embodiment of the invention the peptide of the invention is fused to an epitope tag which provides an epitope to which an anti-tag substance can selectively bind. The epitope tag is generally placed at the amino- or carboxyl-terminus of the peptide but may be incorporated as an internal insertion or substitution as the biological activity permits. The presence of such epitope-tagged forms of a peptide can be detected using a substance such as an antibody against the tagged peptide. Also, provision of the epitope tag enables the peptide to be readily purified by affinity purification using an anti-tag antibody or another type of affinity matrix that binds to the epitope tag. Various tag polypeptides and their respective antibodies are well known in the art. Examples include poly-histidine (poly-his), poly-histidine-glycine (poly-his-gly) tags, the HA tag polypeptide, the c-myc tag, the Strep tag and the FLAG tag.

Fusions also may include the peptides or antigens of this invention fused or coupled to moieties other than amino acids, including lipids and carbohydrates. Further, antigens of this invention may be employed in combination with other vaccinal agents described by the prior art, as well as with other species of vaccinal agents derived from other

microorganisms. Such proteins are useful in the prevention, treatment and diagnosis of diseases caused by a wide spectrum of Streptococcus isolates.

These fusion proteins are constructed for use in the methods and compositions of this invention. These fusion proteins or multimeric proteins may be produced recombinantly, or may be synthesized chemically.

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The peptides of the invention may be prepared by any of a number of conventional techniques. Desired peptides may be chemically synthesized. An alternative approach involves generating the fragments of known peptides by enzymatic digestion, e.g., by treating the protein with an enzyme known to cleave proteins at sites defined by particular amino acid residues, or by digesting the DNA with suitable restriction enzymes, expressing the digested DNA and isolating the desired fragment. Yet another suitable technique involves isolating and amplifying a DNA fragment encoding a desired peptide fragment, by polymerase chain reaction (PCR). Oligonucleotides that define the desired termini of the DNA fragment are employed as the 5' and 3' primers in the PCR. Techniques for making mutations, such as deletions, insertions and substitutions, at predetermined sites in DNA, and therefore in proteins, having a known sequence are well known. One of skill in the art using conventional techniques, such as PCR, may readily use the antigens and peptides provided herein to identify and isolate other similar proteins. Such methods are routine and not considered to require undue experimentation, given the information provided herein. For example, variations can be made using oligonucleotide-mediated sitedirected mutagenesis (Carter et al., Nucl. Acids Res., 13: 4431 (1985); Zoller et al., Nucl. Acids Res. 10: 6487 (1987)), cassette mutagenesis (Wells et al., Gene, 34: 315 (1985)), restriction selection mutagenesis (Wells et al., Philos. Trans. R. Soc. London SerA, 317: 415 (1986)), PCR mutagenesis, or other known techniques can be performed on the cloned DNA to produce the peptide of the invention.

Another subject of the present invention relates to a nucleic acid encoding a peptide of the invention, i.e. any peptide as defined above, or a nucleic acid complementary thereto. Nucleic acid molecules of the present invention may be in the form of RNA, such as mRNA or cRNA, or in the form of DNA, including, for instance, cDNA and genomic DNA e.g. obtained by cloning or produced by chemical synthetic techniques or by a

combination thereof. The DNA may be double- stranded or single-stranded. Single-stranded DNA may be the coding strand, also known as the sense strand, or it may be the non-coding strand, also referred to as the anti-sense strand. Nucleic acid molecule as used herein also refers to, among other, single- and double- stranded DNA, DNA that is a mixture of single- and double-stranded RNA, and RNA that is a mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded, or a mixture of single- and double-stranded regions.

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The nucleic acid may be a fragment of a nucleic acid occurring naturally in *S. pyogenes*, especially in *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, particularly *S. pyogenes* SF370. Preferably the nucleic acid has a sequence as defined in any of the sequences of SEQ ID NOS: 11 to 17 or of any of the homologous variants identified in the attached listing of nucleic acid sequence data. Examples of homologous sequences of a different serotype are those listed below:

Full length nucleic acid sequence (SEQ ID NO)	Nucleic acid of the invention (SEQ ID NO)	Homologous nucleic acid sequences (SEQ ID NOS)
133	11	134 to 143
144	12	145 to 154
144	13	155 to 164
165	14	166 to 175
165	15	176 to 185
165	16	186 to 195
196	17	197 to 206

The nucleic acid also includes sequences that are a result of the degeneration of the genetic code. There are 20 natural amino acids, most of which are specified by more than one codon. Therefore, all nucleotide sequences are included in the invention which result in the peptide as defined above.

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Additionally, the nucleic acid may contain one or more modified bases. Such nucleic acids may also contain modifications e.g. in the ribose-phosphate backbone to increase stability and half life of such molecules in physiological environments. Thus, DNAs or RNAs with backbones modified for stability or for other reasons are "nucleic acid molecule" as that feature is intended herein. Moreover, DNAs or RNAs comprising unusual bases, such as inosine, or modified bases, such as tritylated bases, to name just two examples, are nucleic acid molecule within the context of the present invention. It will be appreciated that a great variety of modifications have been made to DNA and RNA that serve many useful purposes known to those of skill in the art. The term nucleic acid molecule as it is employed herein embraces such chemically, enzymatically or metabolically modified forms of nucleic acid molecule, as well as the chemical forms of DNA and RNA characteristic of viruses and cells, including simple and complex cells, inter alia. For example, nucleotide substitutions can be made which do not affect the polypeptide encoded by the nucleic acid, and thus any nucleic acid molecule which encodes an antigen or fragment or functional active variant thereof as defined above is encompassed by the present invention.

Furthermore, any of the nucleic acid molecules encoding an antigen of the invention or fragment or functional active variant thereof can be functionally linked, using standard techniques such as standard cloning techniques, to any desired regulatory sequences, whether a *S. pyogenes* regulatory sequence or a heterologous regulatory sequence, heterologous leader sequence, heterologous marker sequence or a heterologous coding sequence to create a fusion protein.

- The nucleic acid of the invention may be originally formed *in vitro* or in a cell in culture, in general, by the manipulation of nucleic acids by endonucleases and/or exonucleases and/or polymerases and/or ligases and/or recombinases or other methods known to the skilled practitioner to produce the nucleic acids.
- In one embodiment of the invention the nucleic acid is located in a vector. A vector may additionally include nucleic acid sequences that permit it to replicate in the host cell, such as an origin of replication, one or more desired genes and/or selectable marker genes and other genetic elements known in the art such as regulatory elements directing transcription,

translation and/or secretion of the encoded protein. The vector may be used to transduce, transform or infect a cell, thereby causing the cell to express inserted nucleic acids and/or proteins other than those native to the cell. The vector optionally includes materials to aid in achieving entry of the nucleic acid into the cell, such as a viral particle, liposome, protein coating or the like. Numerous types of appropriate expression vectors are known in the art for protein expression, by standard molecular biology techniques. Such vectors are selected from among conventional vector types including insects, e.g., baculovirus expression, or yeast, fungal, bacterial or viral expression systems. Other appropriate expression vectors, of which numerous types are known in the art, can also be used for this purpose. Methods for obtaining such expression vectors are well-known (see, e.g. Sambrook et al, Molecular Cloning. A Laboratory Manual, 2nd edition, Cold Spring Harbor Laboratory, New York (1989)). In one embodiment, the vector is a viral vector. Viral vectors include, but are not limited to, retroviral and adenoviral vectors.

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Suitable host cells or cell lines for transfection by this method include bacterial cells. For example, the various strains of *E. coli* are well-known as host cells in the field of biotechnology. Various strains of *B. subtilis*, *Pseudomonas*, *Streptomyces*, and other bacilli and the like may also be employed in this method. Many strains of yeast cells known to those skilled in the art are also available as host cells for expression of the peptides of the present invention. Other fungal cells or insect cells such as *Spodoptera frugipedera* (Sf9) cells may also be employed as expression systems. Alternatively, mammalian cells, such as human 293 cells, Chinese hamster ovary cells (CHO), the monkey COS-1 cell line or murine 3T3 cells derived from Swiss, BALB/c or NIH mice may be used. Still other suitable host cells, as well as methods for transfection, culture, amplification, screening, production, and purification are known in the art.

A peptide of the invention may be produced by expressing a nucleic acid of the invention in a suitable host cell. The host cells can be transfected, e.g. by conventional means such as electroporation with at least one expression vector containing a nucleic acid of the invention under the control of a transcriptional regulatory sequence. The transfected or transformed host cell is then cultured under conditions that allow expression of the protein. The expressed protein is recovered, isolated, and optionally purified from the cell (or from the culture medium, if expressed extracellularly) by appropriate means known to one of

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skill in the art. For example, the proteins are isolated in soluble form following cell lysis, or extracted using known techniques, e.g. in guanidine chloride. If desired, the peptides or fragments of the invention are produced as a fusion protein. Such fusion proteins are those described above. Alternatively, for example, it may be desirable to produce fusion proteins to enhance expression of the protein in a selected host cell or to improve purification. The molecules comprising the peptides and antigens of this invention may be further purified using any of a variety of conventional methods including, but not limited to: liquid chromatography such as normal or reversed phase, using HPLC, FPLC and the like; affinity chromatography (such as with inorganic ligands or monoclonal antibodies); size exclusion chromatography; immobilized metal chelate chromatography; gel electrophoresis; and the like. One of skill in the art may select the most appropriate isolation and purification techniques without departing from the scope of this invention. Such purification provides the antigen in a form substantially free from other proteinaceous and non-proteinaceous materials of the microorganism.

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Another subject of the invention is a pharmaceutical composition, especially a vaccine, comprising

- (i) at least one peptide according to the invention, and/or
- (ii) at least one peptide comprising or consisting of the sequence of any of the SEQ ID NO: 8, SEQ ID NO: 9, or SEQ ID NO: 10, or a functional active variant thereof, and
- (iii) optionally a pharmaceutically acceptable carrier or excipient.

The variants of the peptides of (ii) are as defined and may be obtained as the peptides of (i) (see above description of the peptides of the invention). Preferred alterations of the sequences of SEQ ID NO: 8 or 10 are those listed in tables 8 and 9, respectively.

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The peptides of (i) and (ii) are referred to as pharmaceutical peptides of the invention.

With respect to the peptide of (ii), these proteins have been shown for the first time to be capable to provide protection against lethal *S. pyogenes* challenge (see Example 1), particularly in a physiologically highly relevant intranasal challenge model. Especially protein Spy0895 (SEQ ID NO: 9) shows particular promise as a vaccine candidate, because it provided protection against group A streptococcal infection in all three models listed in Table 1.

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A pharmaceutical peptide of the invention may be used for methods for immunizing or treating humans and/or animals with the disease caused by infection with S. pyogenes. Therefore, the pharmaceutical peptide may be used within a pharmaceutical composition. The pharmaceutical composition of the present invention may further encompass pharmaceutically acceptable carriers and/or excipients. The pharmaceutically acceptable carriers and/or excipients useful in this invention are conventional and may include buffers, stabilizers, diluents, preservatives, and solubilizers. Remington's Pharmaceutical Sciences, by E. W. Martin, Mack Publishing Co., Easton, PA, 15th Edition (1975), describes compositions and formulations suitable for pharmaceutical delivery of the (poly)peptides herein disclosed. In general, the nature of the carrier or excipients will depend on the particular mode of administration being employed. For instance, parenteral formulations usually comprise injectable fluids that include pharmaceutically and physiologically acceptable fluids such as water, physiological saline, balanced salt solutions, aqueous dextrose, glycerol or the like as a vehicle. For solid compositions (e. g. powder, pill, tablet, or capsule forms), conventional non-toxic solid carriers can include, for example, pharmaceutical grades of mannitol, lactose, starch, or magnesium stearate. In addition to biologically neutral carriers, pharmaceutical compositions to be administered can contain minor amounts of non-toxic auxiliary substances, such as wetting or emulsifying agents, preservatives, and pH buffering agents and the like, for example sodium acetate or sorbitan monolaurate.

In a preferred embodiment the pharmaceutical composition further comprises an immunostimulatory substance such as an adjuvant. The adjuvant can be selected based on the method of administration and may include mineral oil-based adjuvants such as Freund's complete and incomplete adjuvant, Montanide incomplete Seppic adjuvant such as ISA, oil in water emulsion adjuvants such as the Ribi adjuvant system, syntax adjuvant formulation containing muramyl dipeptide, IC31TM (Intercell; a synthetic adjuvant comprising the peptide motif KLK [WO 02/32451] and an oligonucleotide [WO 01/93905]), or aluminum salt adjuvants. Preferably, the adjuvant is a mineral oil-based adjuvant, most preferably ISA206 (SEPPIC, Paris, France).

In other embodiments the immunostimulatory substance is selected from the group comprising polycationic polymers, especially polycationic peptides such as polyarginine, immunostimulatory deoxynucleotides (ODNs), especially Oligo(dIdC)₁₃, peptides containing at least two LysLeuLys motifs, especially KLKLLLLKLK (SEQ ID NO: 55), neuroactive compounds, especially human growth hormone, alum, adjuvants or combinations thereof. In further embodiments, the combination is either a polycationic polymer and immunostimulatory deoxynucleotides or of a peptide containing at least two

LysLeuLys motifs and immunostimulatory deoxynucleotides. In a still another

embodiment the polycationic polymer is a polycationic peptide.

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In an embodiment the immunostimulatory substance is at least one immunostimulatory nucleic acid. Immunostimulatory nucleic acids are e.g. neutral or artificial CpG containing nucleic acids, short stretches of nucleic acids derived from non-vertebrates or in form of short oligonucleotides (ODNs) containing non-methylated cytosine-guanine dinucleotides (CpG) in a defined base context (e.g. as described in WO 96/02555). Alternatively, also nucleic acids based on inosine and cytidine as e.g. described in WO 01/93903, or deoxynucleic acids containing deoxy-inosine and/or deoxyuridine residues (described in WO 01/93905 and WO 02/095027) may preferably be used as immunostimulatory nucleic acids in the present invention. Preferably, mixtures of different immunostimulatory nucleic acids are used in the present invention. Additionally, the aforementioned polycationic compounds may be combined with any of the immunostimulatory nucleic acids as aforementioned. Preferably, such combinations are according to the ones described in WO 01/93905, WO 02/32451, WO 01/54720, WO 01/93903, WO 02/13857, WO 02/095027 and WO 03/047602.

In addition or alternatively, such pharmaceutical or vaccine composition may comprise a neuroactive compound. Preferably, the neuroactive compound is human growth factor, e.g. described in WO 01/24822. Also preferably, the neuroactive compound is combined with any of the polycationic compounds and/or immunostimulatory nucleic acids as defined above.

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The composition may be used e.g. for immunization or treatment of a subject. The pharmaceutical composition encompasses at least one pharmaceutical peptide of the invention; however, it may also contain a cocktail (i.e., a simple mixture) containing different pharmaceutical peptides (including fragments and other variants) of the invention, optionally mixed with different antigenic proteins or peptides of other pathogens. Such mixtures of these peptides, polypeptides, proteins or fragments or variants thereof are useful e.g. in the generation of desired antibodies to a wide spectrum of Streptococci isolates. The pharmaceutical peptide(s) of the present invention may also be used in the form of a pharmaceutically acceptable salt. Suitable acids and bases which are capable of forming salts with the peptides of the present invention are well known to those of skill in the art, and include inorganic and organic acids and bases.

Still another subject of the invention is a pharmaceutical composition containing a nucleic acid selected from the group consisting of:

- (i) a nucleic acid of the invention and/or a nucleic acid complementary thereto, and/or
- (ii) a nucleic acid coding for the peptide comprising or consisting of the sequence of any of the SEQ ID NO: 8, SEQ ID NO: 9, or SEQ ID NO: 10, particularly a DNA sequence of any of the SEQ ID NO: 18, SEQ ID NO: 19, or SEQ ID NO: 20, or a functional active variant thereof or a nucleic acid complementary thereto or the corresponding RNA sequence, and
- (iii) optionally a pharmaceutically acceptable carrier or excipient.

The variants of the nucleic acids of (ii) are as defined and may be obtained as the nucleic acids of (i) (see above description of the nucleic acids of the invention). The nucleic acids of (i) and (ii) are referred to as pharmaceutical nucleic acids of the invention.

The pharmaceutical nucleic acid sequences, alone or in combination with other nucleic acid sequences encoding antigens or antibodies or directed to other pathogenic microorganisms, may further be used as components of a pharmaceutical composition. The composition may be used for immunizing or treating humans and/or animals being susceptible to or having a disease caused by infection with *S. pyogenes*, particularly *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370. The pharmaceutically acceptable carrier or excipient may be as defined above.

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In another embodiment, the pharmaceutical nucleic acids of this invention, alone or in combination with nucleic acid sequences encoding other antigens or antibodies from other pathogenic microorganisms, may further be used in compositions directed to actively induce a protective immune response in a subject to the pathogen. These components of the present invention are useful in methods for inducing a protective immune response in humans and/or animals against infection with *S. pyogenes*, particularly with *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370.

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For use in the preparation of the therapeutic or vaccine compositions, nucleic acid delivery compositions and methods are useful, which are known to those of skill in the art. The pharmaceutical nucleic acid of the invention may be employed in the methods of this invention or in the compositions described herein as DNA sequences, either administered as naked DNA, or associated with a pharmaceutically acceptable carrier and provide for *in vivo* expression of the antigen, peptide or polypeptide. So-called "naked DNA" may be used to express the antigen, peptide or polypeptide of the invention *in vivo* in a patient. (See, e.g., J. Cohen, Science, 259: 1691-1692, which describes similar uses of "naked DNA"). For example, "naked DNA" associated with regulatory sequences may be administered therapeutically or as part of the vaccine composition e.g., by injection.

Alternatively, a nucleic acid, especially a pharmaceutical nucleic acid according to the invention, encoding an antigen or peptide of the invention or a nucleic acid complementary

thereto may be used within a pharmaceutical composition, e.g. in order to express the antigen or (pharmaceutical) peptide of the invention in vivo, e.g., to induce antibodies.

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A preferred embodiment of the invention relates to a pharmaceutical composition, wherein the pharmaceutical nucleic acid according to the invention is comprised in a vector and/or a cell. Vectors and cells suitable in the context of the present invention are described above. Vectors are particularly employed for a DNA vaccine. An appropriate vector for delivery may be readily selected by one of skill in the art. Exemplary vectors for *in vivo* gene delivery are readily available from a variety of academic and commercial sources, and include, e.g., adeno-associated virus (International patent application No. PCT/US91/03440), adenovirus vectors (M. Kay et al, Proc. Natl. Acad. Sci. USA, 91: 2353 (1994); S. Ishibashi et al, J. Clin. Invest., 92: 883 (1993)), or other viral vectors, e.g., various poxviruses, vaccinia, etc.. Recombinant viral vectors, such as retroviruses or adenoviruses, are preferred for integrating the exogenous DNA into the chromosome of the cell.

Another subject of the invention relates to an antibody or functional active fragment thereof which binds specifically to the antigen of the invention. The present invention includes, for example, monoclonal and polyclonal antibodies, chimeric, single chain, and humanized antibodies, as well as Fab fragments, or the product of a Fab expression library.

While *S. pyogenes* infections are primarily a disease of children and cause non-severe diseases such as bacterial pharyngitis and impetigo, GAS are also responsible for streptococcal toxic shock syndrome associated necrotizing fasciitis (Cone, L., et al. (1987). New Engl J Med 317: 146-9; Stevens, D. (1992). Clin Infect Dis 14: 2-11) and several post-streptococcal sequelae such as acute rheumatic fever, acute glomerulonephritis and reactive arthritis. It would be very beneficial to provide monoclonal or polyclonal antibody therapies which target antigenic proteins of *S. pyogenes* and have the potential to support a therapy of an infection or eliminate the pathogen and the disease altogether.

In a preferred embodiment the antibody is a monoclonal, polyclonal, chimeric or humanized antibody or functional active variant thereof. In another preferred embodiment the functional active fragment comprises a Fab fragment.

Antibodies generated against the antigens, fragments or variants thereof of the present invention can be obtained by direct injection of the antigens, fragments or variants thereof into an animal or by administering the antigens, fragments or variants thereof to an animal, preferably a non-human. The antibody so obtained will then bind the antigens, fragments or variants. Such antibodies can then be used to isolate reactive antigens, fragments or variants thereof from tissue expressing those.

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For preparation of monoclonal antibodies, any technique known in the art, which provides antibodies produced by continuous cell line cultures, e.g. a hybridoma cell line, can be used.

Techniques described for the production of single chain antibodies (U. S. Patent No. 4,946,778) can be adapted to produce single chain antibodies to the antigens, fragments or variants thereof according to this invention. Also, transgenic mice or other organisms such as other mammals may be used to express humanized antibodies to antigens, fragments or variants thereof according to this invention.

Still another subject of the invention relates to a hybridoma cell line which produces the antibody of the invention.

Hybridoma cell lines expressing desirable monoclonal antibodies are generated by well-known conventional techniques. The hybridoma cell can be generated by fusing a normal-activated, antibody-producing B cell with a myeloma cell. In the context of the present invention the hybridoma cell is able to produce an antibody specifically binding to the antigen of the invention.

Similarly, desirable high titre antibodies are generated by applying known recombinant techniques to the monoclonal or polyclonal antibodies developed to these antigens (see, e.g., PCT Patent Application No. PCT/GB85/00392; British Patent Application Publication No. GB2188638A; Amit et al., Science, 233: 747-753 (1986); Queen et al., Proc. Natl. Acad. Sci. USA, 86: 10029-10033 (1989); PCT Patent Application No. WO90/07861;

Riechmann et al., Nature, 332: 323-327 (1988); Huse et al., Science, 246: 1275-1281 (1988)).

The present invention also provides a method for producing an antibody according to the invention, characterized by the following steps:

- (a) administering an effective amount of the peptide according to the invention to an animal; and
- (b) isolating the antibody produced by the animal in response to the administration of step (a) from the animal.

Another subject of the invention relates to a method for producing an antibody according to the invention, characterized by the following steps:

- (a) contacting a B cell with an effective amount of the peptide according to the invention;
- (b) fusing the B cell of step (a) with a myeloma cell to obtain a hybridoma cell; and
- (c) isolating the antibody produced by the cultivated hybridoma cell.

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More particularly, the antibody may be produced by initiating an immune response in a non-human animal by administrating a peptide of the invention to an animal, removing an antibody containing body fluid from said animal, and producing the antibody by subjecting said antibody containing body fluid to further purification steps. Alternatively, the antibody may be produced by initiating an immune response in a non-human animal by administrating an antigen, fragment or variant thereof, as defined in the present invention, to said animal, removing the spleen or spleen cells from said animal and/or producing hybridoma cells of said spleen or spleen cells, selecting and cloning hybridoma cells specific for said antigen, fragment or variant thereof and producing the antibody by cultivation of said cloned hybridoma cells.

In a preferred embodiment the antibody produced according to a method of the invention is additionally purified. Methods of purification are known to the skilled artisan.

The antibody may be used in methods for preventing or treating an infection. Accordingly, still another subject of the invention relates to a pharmaceutical composition, especially a

vaccine, comprising an antibody of the invention. The pharmaceutical composition may encompass further components as detailed above. The composition may further encompass substances increasing their capacity to stimulate T cells. These include T helper cell epitopes, lipids or liposomes or preferred modifications as described in WO01/78767. Another way to increase the T cell stimulating capacity of epitopes is their formulation with immune stimulating substances for instance cytokines or chemokines like interleukin-2, -7, -12, -18, class I and II interferons (IFN), especially IFN-gamma, GM-CSF, TNF-alpha, flt3-ligand and others.

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A further subject of the invention relates to a pharmaceutical composition comprising the pharmaceutical peptide of the invention or the pharmaceutical nucleic acid of the invention or an antibody of the invention or functional fragment thereof for the immunization of a subject against an infection or the treatment of a subject having an infection, wherein the infection is preferably a S. pyogenes infection. In another aspect of the invention a pharmaceutical peptide of the invention or a pharmaceutical nucleic acid of the invention or an antibody of the invention or functional fragment thereof is used for the manufacture of a medicament for the immunization of a subject against an infection or the treatment of a subject having an infection, wherein the infection is preferably a S. pyogenes infection, more preferably an infection with S. pyogenes serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially S. pyogenes SF370. Alternatively, a pharmaceutical peptide or a pharmaceutical nucleic acid of the invention or an antibody of the invention or functional fragment thereof is used in a method of immunizing or treating a subject in need thereof, wherein an effective amount of the pharmaceutical peptide or the pharmaceutical nucleic acid of the invention or an antibody of the invention or functional fragment thereof is administered to the subject. The subject may be immunized in order to prevent an infection, particularly a S. pyogenes infection, or may be treated to ameliorate or cure an infection, particularly a S. pyogenes infection. The determination of the effective amount to be administered is within the knowledge of the skilled practitioner. Exemplary amounts are mentioned below.

The pharmaceutical peptides or the pharmaceutical nucleic acids of the invention are generally useful for inducing an immune response in a subject. The vaccine used for

immunization may be administered to a subject susceptible to infection by S. pyogenes, preferably mammals, and still more preferably humans. Potential modes of administration include oral, intranasal, intramuscular, intra-lymph node, intradermal, intraperitoneal, subcutaneous, and combinations thereof, but most preferably intramuscular injection. The volume of the dose for intramuscular administration is preferably up to about 5 mL, for example, between 0.3 mL and 3 mL, between 1 mL and 3 mL, about 0.5 to 1 mL, or about 2 mL. The amount of protein comprising the antigen in each dose should be enough to confer effective immunity to decrease the risk of developing clinical signs, e.g. resulting from S. pyogenes infection. In different embodiments, the unit dose of protein should be up to about 5 µg protein/kg body weight, between about 0.2 to 3 µg, between about 0.3 to 1.5 μg , between about 0.4 to 0.8 μg , or about 0.6 μg . In alternative embodiments unit doses of protein could be up to about 6 µg protein/kg body weight, between about 0.05 to 5 µg, or between about 0.1 to 4 μg . In different embodiments, the dose is administered 1 to 3 times, e.g. with an interval of 1 to 3 weeks. Representative amounts of protein per dose are from approximately 1 μg to approximately 1 mg, more preferably from approximately 5 μg to approximately 500 µg, still more preferably from approximately 10 µg to approximately $250~\mu g$ and most preferably from approximately $25~\mu g$ to approximately $100~\mu g$.

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In still another aspect of the invention the antibody of the invention or functional fragment thereof is used for the manufacture of a medicament for the treatment of an infection, preferably a *S. pyogenes* infection, more preferably an infection with *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370. Alternatively, the antibody of the invention is used in a method of treating a subject in need thereof, wherein an effective amount of the antibody of the invention is administered to the subject. The subject may be treated to ameliorate or cure an infection, particularly a *S. pyogenes* infection. The determination of the effective amount to be administered is within the knowledge of the skilled practitioner.

The treatment involves administering an effective amount of an antibody of the invention to a subject, preferably a mammal, more preferably a human. Thus, antibodies against the antigens, fragments or variants thereof of the present invention may be employed to inhibit and/or treat infections, particularly bacterial infections and especially infections arising

from *S. pyogenes*, especially *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370.

An "effective amount" of a pharmaceutical peptide, a pharmaceutical nucleic acid or an antibody of the invention may be calculated as that amount capable of exhibiting an *in vivo* effect, e.g. preventing or ameliorating a sign or symptom of infection, particularly *S. pyogenes* infection, especially of *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370. Such amounts may be determined by one of skill in the art. Preferably, such a composition is administered parenterally, preferably intramuscularly or subcutaneously. However, it may also be formulated to be administered by any other suitable route, including orally or topically. The selection of the route of delivery and dosage of such therapeutic compositions is within the skill of the art.

Treatment in the context of the present invention refers to both therapeutic treatment and prophylactic or preventative measures, wherein the object is to prevent or slow down (lessen) the targeted pathologic condition or disorder. Those in need of treatment include those already with the disorder as well as those prone to have the disorder or those in whom the disorder is to be prevented.

Another subject of the invention relates to a method of diagnosing a *S. pyogenes* infection comprising the steps of:

- 25 (a) contacting a sample obtained from a subject with the peptide according to the invention; and
 - (b) detecting the presence of an antibody against S. pyogenes in the sample.

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The peptides of the invention may be used for the detection of the *S. pyogenes*, particularly *S. pyogenes* serotype M1, M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118, especially *S. pyogenes* SF370. Preferably such detection is for diagnosis, more preferable for the diagnosis of a disease, most preferably for the diagnosis of a *S. pyogenes* infection. The

peptides or polypeptides may be used to detect the presence of a *S. pyogenes*-specific antibody or fragment thereof e.g. in a sample obtained from a subject. The sample may be e.g. a blood sample. Alternatively, the presence of a *S. pyogenes*-specific antigen can be

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Accordingly, an alternative method of diagnosing a *S. pyogenes* infection comprises the steps of:

- (a) contacting a sample obtained from a subject with the antibody according to the invention; and
- (b) detecting the presence of an antigen of S. pyogenes in the sample.

detected using an antibody of the invention.

The present invention also relates to diagnostic assays such as quantitative and diagnostic assays for detecting levels of the peptides or antibodies of the present invention in cells and tissues or body fluids, including determination of normal and abnormal levels. Assay techniques that can be used to determine levels of a peptide or an antibody, in a sample derived from a host are well known to those of skill in the art. Such assay methods include radioimmunoassays, competitive-binding assays, Western Blot analysis and ELISA assays. Among these, ELISAs frequently are preferred. An ELISA assay initially comprises preparing an antibody specific to the peptide, particularly the antigen, preferably a monoclonal antibody. In addition, a reporter antibody generally is prepared which binds to the monoclonal antibody. The reporter antibody is attached to a detectable reagent such as radioactive, fluorescent or enzymatic reagent, such as horseradish peroxidase enzyme.

The peptides or antibodies of the present invention may also be used for the purpose of or in connection with an array. More particularly, at least one of the peptides or antibodies of the present invention may be immobilized on a support. Said support typically comprises a variety of antigens and fragments thereof whereby the variety may be created by using one or several of the peptides or antibodies of the present invention. The characterizing feature of such array as well as of any array in general is the fact that at a distinct or predefined region or position on said support or a surface thereof, a distinct polypeptide is immobilized. Because of this any activity at a distinct position or region of an array can be correlated with a specific polypeptide. The number of different peptides or antibodies of

the present invention immobilized on a support may range from as little as 10 to several 1000 different peptides or antibodies of the present invention.

The manufacture of such arrays is known to the one skilled in the art and, for example, described in US patent 5,744,309. The array preferably comprises a planar, porous or non-porous solid support having at least a first surface. Preferred support materials are, among others, glass or cellulose. It is also within the present invention that the array is used for any of the diagnostic applications described herein. Apart from the peptides or antibodies of the present invention also the nucleic acid molecules according to the present invention may be used for the generation of an array as described above.

Another aspect of the invention relates to a method for identifying a ligand capable of binding to a peptide according to the invention comprising:

(a) providing a test system comprising the peptide,

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- (b) contacting the test system with a test compound, and
 - (c) detecting a signal generated in response to the binding of the test compound to the peptide.

More particularly, the method may be carried out by contacting an isolated or immobilized peptide according to the invention with a candidate ligand under conditions to permit binding of the candidate ligand to the peptide, wherein the test system comprises a component capable of providing a detectable signal in response to the binding of the candidate ligand to said peptide; and detecting the presence or absence of a signal generated in response to the binding of the ligand to the peptide. The ligand may be an agonist or an antagonist.

Test systems for detection binding of a ligand are known to the skilled artisan and include e.g. binding assays with labeled ligand such as radioligands, fluorescence-labeled ligands or enzyme-labeled ligands.

The test compound can be any test compound either naturally occurring or chemically synthesized. Naturally occurring test compounds include in particular antibodies, preferably those showing similarity to the antibodies of the invention. In one preferred

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embodiment of the invention the test compound is provided in the form of a chemical compound library. Chemical compound libraries include a plurality of chemical compounds and have been assembled from any of multiple sources, including chemically synthesized molecules and natural products, or have been generated by combinatorial chemistry techniques. They are especially suitable for high throughput screening. They may be comprised of chemical compounds of a particular structure or compounds of a particular creature such as a plant.

The method for identifying a ligand may also include the following steps:

- (a) providing a peptide according to the invention,
 - (b) providing an interaction partner to the peptide especially an antibody according to the invention,
 - (c) allowing interaction of the peptide to said interaction partner to form a interaction complex,
- 15 (d) providing a test compound,
 - (e) allowing a competition reaction to occur between the test compound and the interaction complex, and
 - (f) determining whether the test compound inhibits or reduces the interaction activities of the peptide with the interaction partner.

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The ligands identified may be employed, for instance, to inhibit diseases arising from infection with Streptococcus, especially *S. pyogenes* and may therefore be formulated in a pharmaceutical composition.

In a last aspect, the peptide according to the invention is used for the isolation and/or purification and/or identification of a ligand of the peptide, wherein the isolation and/or purification and/or identification of the ligand may be carried out as detailed above or as known to the person skilled in the art. In a preferred embodiment of the invention an affinity device may be used. The affinity device may comprise as least a support material and any peptide according to the present invention, which is attached to the support material. Because of the specificity of the peptides according to the present invention for their target cells or target molecules or their interaction partners, the peptides allow a selective removal of their interaction partner(s) from any kind of sample applied to the

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support material provided that the conditions for binding are met. The sample may be a biological or medical sample, including but not limited to, fermentation broth, cell debris, cell preparation, tissue preparation, organ preparation, blood, urine, lymph liquid, liquor and the like. The peptide may be attached to the matrix in a covalent or non-covalent manner. Suitable support material is known to the one skilled in the art and can be selected from the group comprising cellulose, silicon, glass, aluminium, paramagnetic beads, starch and dextrane.

The present invention is further illustrated by the following figures, examples and the sequence data, from which further features, embodiments and advantages may be taken. It is to be understood that the present examples are given by way of illustration only and not by way of limitation of the disclosure.

Figure 1 shows the protection achieved by active immunization with selected *S. pyogenes* antigens and sub-constructs in a mouse lethality model.

Figure 2 shows the protection achieved by active immunization with selected *S. pyogenes* antigens and sub-constructs in a mouse lethality model.

Figure 3 shows the protection achieved by active immunization with selected *S. pyogenes* antigens and sub-constructs in a mouse lethality model.

Figure 4 shows the protection achieved by active immunization with selected *S. pyogenes* antigens in a mouse lethality model.

Table 1 shows the recombinant proteins of *S. pyogenes* and fragments thereof assessed for protection in murine models of infection.

Table 2 shows the oligonucleotides used for the cloning of genes encoding antigenic proteins and fragments thereof of *S. pyogenes*.

Table 3 shows the *S. pyogenes* strains used for the gene conservation study.

Table 4 shows the oligonucleotides used for PCR and sequencing of the *S. pyogenes* genes.

Table 5 shows the variable amino acid positions of Spy0269 from S. pyogenes strains.

Table 6 shows the variable amino acid positions of Spy0292 from S. pyogenes strains.

Table 7 shows the variable amino acid positions of Spy0416 from S. pyogenes strains.

Table 8 shows the variable amino acid positions of Spy0488 from S. pyogenes strains.

Table 9 shows the variable amino acid positions of Spy0872 from S. pyogenes strains.

Table 10 shows the variable amino acid positions of Spy0895 from S. pyogenes strains.

Table 11 shows the variable amino acid positions of Spy1536 from S. pyogenes strains.

Table 12 shows the variable amino acid positions of Spy1666 from S. pyogenes strains.

FIGURES

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Figure 1: Protection achieved by active immunization with selected S. pyogenes antigens and sub-constructs in a mouse lethality model. CD-1 mice (10 mice per group) were immunized subcutaneously with recombinant antigens cloned from an emm type 1 S. pyogenes strain (SF370) and challenged with the MA-A20 (emm type 23) strain. Survival was monitored for 14 days post-challenge. Mice were immunized subcutaneously with 50 µg recombinant protein adjuvanted with CFA/IFA. (A) Spy0292, and its sub-constructs Spy0292-1 and Spy0292-3; Spy0488; (B) Spy0872 and its sub-construct Spy0872-2. Anesthetized mice were challenged intranasally with 10⁸ cfu S. pyogenes MA-A20. Adjuvant control mice were used as negative controls, while M1 (Spy2018) served as positive control. Numbers of surviving mice are plotted as percentage of total mice.

Figure 2: Protection achieved by active immunization with selected S. pyogenes antigens and sub-constructs in a mouse lethality model. CD-1 mice (10 mice per group) were immunized subcutaneously with recombinant antigens cloned from an emm type 1 S. pyogenes strain (SF370) and challenged with the MA-A20 (emm type 23) strain. Survival was monitored for 14 days post-challenge. Mice were immunized subcutaneously with 50 µg recombinant protein adjuvanted with CFA/IFA. (A) Spy0269 and its sub-construct Spy0269-1; (B) Spy0416A and 3 sub-constructs (Spy0416A-1, Spy0416A-6 and Spy0416A-7) and Spy0416B. Anesthetized mice were challenged intranasally with 10⁸ cfu S. pyogenes MA-A20. Adjuvant control mice were used as negative controls, while M1 protein (Spy2018) served as positive control. Numbers of surviving mice are plotted as percentage of total mice.

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- Figure 3: Protection achieved by active immunization with selected S. pyogenes antigens or sub-constructs in a mouse lethality model. CD-1 mice (10 mice per group) were immunized subcutaneously with recombinant antigens cloned from an emm type 1 S. pyogenes strain (SF370) and challenged with the MA-A20 (emm type 23) strain. Survival was monitored for 14 days post-challenge. Mice were immunized subcutaneously with 50 μg recombinant protein adjuvanted with aluminum hydroxide. (A) Spy1727, Spy0269-1, Spy0872-2, and Spy0416A-1; (B) Spy1666, Spy1536, Spy0895, and Spy0292-1. Anesthetized mice were challenged intranasally with 10⁸ cfu S. pyogenes MA-A20. Adjuvant control mice were used as negative controls, while M1 protein (Spy2018) served as positive control. Numbers of surviving mice are plotted as percentage of total mice.
- **Figure 4: Protection achieved by active immunization with selected** *S. pyogenes* **antigens in a mouse lethality model.** BALB/c mice (10 mice per group) were immunized intranasally with recombinant antigens cloned from an emm type 1 *S. pyogenes* strain (SF370) and challenged either with **(A)** MA-A20 (emm type 23) strain or with **(B)** MA-A147 (emm type 11/106) strain. Survival was monitored for 14 days post-challenge. Mice were immunized intranasally with 30-50 μg recombinant protein adjuvanted with IC31TM. **(A)** Spy1536 and Spy0895; **(B)** Spy1727 and Spy1536. Anesthetized mice were challenged intranasally with 10⁶ cfu *S. pyogenes* MA-A20 or 10⁸ cfu *S. pyogenes* MA-A147. Adjuvant control mice were used as negative controls, while M1 protein (Spy2018) served as positive control. Numbers of surviving mice are plotted as percentage of total mice.

EXAMPLES

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Example 1: Group A streptococcal antigens and fragments thereof inducing protective immune responses against lethal sepsis in intranasal challenge models.

Experimental procedures

Cloning and expression of recombinant pneumococcal proteins

Cloning of genes / DNA fragments:

The gene/DNA fragment of interest (see Table 1) was amplified from genomic DNA of *Streptococcus pyogenes* SF370 (serotype M1) by PCR using gene specific primers (see Table 2). Apart from the gene specific part, the primers had restriction sites that aided in a directional cloning of the amplified PCR product. The gene annealing (specific) part of the primer ranged between 15-30 bases in length. The PCR products obtained were digested with the appropriate restriction enzymes and cloned into the pET28b (+) vector (Novagen) for His-tagged proteins. The constructs including full length and fragments of the selected antigens are listed in Table 1. Once the recombinant plasmid was confirmed to contain the gene of interest, *E. coli* BL21 star® cells (Invitrogen) that served as expression host were transformed.

Expression and purification of proteins:

E coli BL21 star® cells harboring the recombinant plasmid were grown into log phase in the required culture volume. Once an OD_{600nm} of 0.6 was reached the culture was induced with 0.5 mM IPTG (isopropyl-beta-D-thiogalactopyranoside) at 37°C for 3 hours. The cells were harvested by centrifugation, lysed by a combination of the freeze-thaw method followed by disruption of cells with BugBuster® (Novagen). The lysate was separated by centrifugation into soluble (supernatant) and insoluble (pellet) fractions. Depending on the location of the protein different purification strategies were applied.

A) If the His-tagged protein was in the soluble fraction, protein purification was done by binding the supernatant to Ni-Sepharose beads (Ni-SepharoseTM 6 Fast Flow, GE Healthcare). Due to the presence of the hexa Histidine (6xHIS) at the C terminus of the

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expressed protein, it bound to the Ni-Sepharose while the other contaminating proteins were washed from the column by wash buffer. The protein was eluted by 500 mM Imidazole in 20 mM NaH₂PO₄, 0.5 mM NaCl buffer at pH 7.4. The eluate was concentrated, assayed by Bradford for protein concentration and checked by SDS-PAGE and Western blot.

B) If the protein was present in the insoluble fraction the pellet was solubilized in suitable buffer containing 8 M urea and applied onto the Ni-NTA column under denaturing conditions (in buffer containing 8 M urea) using the same materials and procedure as mentioned above. Contaminating proteins were washed from the column by wash buffer without urea. Refolding of the His-tagged protein was performed while the protein was immobilized on the Ni-NTA matrix. After renaturation, proteins were eluted by the addition of 500 mM Imidazole. The eluate was dialyzed to remove traces of urea and concentrated if the volume was large, checked by SDS-PAGE and measured by the Bradford method.

Animal protection studies

Animals:

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20 CD-1 or BALB/c female mice (6 - 8 weeks) were used.

Active immunization (subcutaneous route):

50 μg of recombinant proteins buffered in PBS were injected subcutaneously into CD-1 mice (volume 100 μL), adjuvanted with Complete Freund adjuvant (CFA, final concentration: 50%), aluminium hydroxide (ALUM, final concentration: 1%) or IC31TM (final concentration: 100 nmol L-KLKLLLLKLK (SEQ ID NO: 55), 4 nmol oligodexoynucleotide ODN1a (dIdC)₁₃ in PBS) (Intercell AG, Vienna, Austria). Animals were boosted twice with the same amount of protein and adjuvant (except for CFA where Incomplete Freund adjuvant (IFA) was used for the booster immunizations; final concentration: 50%), at days 14 and 28. The published (Dale et al., J. Immunol. 151: 2188 (1993)) protective M1 or M23 protein antigens were used as positive controls, while mice immunized with adjuvant only served as negative controls. Antibody titers were measured at day 35 by ELISA using the respective recombinant proteins.

Active immunization (intranasal route):

30 - 50 µg of recombinant proteins buffered in PBS were injected intranasally into BALB/c mice (volume 20 µL), adjuvanted with IC31TM (final concentration: 10 nmol L-KLKLLLLKLK (SEQ ID NO: 55), 0.4 nmol oligodexoynucleotide ODN1a (dIdC)₁₃ in PBS) (Intercell AG, Vienna, Austria). Animals were boosted three times with the same amount of protein and adjuvant at days 7, 14 and 28. The published protective M1 or M23 protein antigens were used as positive controls, while mice immunized with adjuvant only served as negative controls. Antibody titers were measured at day 35 by ELISA using the respective recombinant proteins.

Bacterial challenge:

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Freshly grown *S. pyogenes* strains MA-A20 or MA-A147 were used. 1 mL bacterial suspension from an o/n culture of the respective *S. pyogenes* strain was added to 50 mL THY culture medium. Optical density was measured until the bacterial suspension reached an OD_{600nm} between 0.4 and 0.6. Bacterial counts were determined using an individually established growth curve. Bacterial cells were spun down and adjusted with PBS to obtain the desired cfu count. In order to determine the viable cell numbers present in the bacterial inoculum, cfus were determined via plating on blood agar plates. 10^6 - 10^8 cfus were applied intranasally (20 μ L) into individually anesthetized mice. Protection by immunization was measured by a bacteraemia / sepsis model where survival rates were followed for 2 to 3 weeks post-challenge and survival was expressed in percentage of the total number of animals (10 mice / group).

25 Results

Group A streptococcal antigens and/or their fragments were identified showing protection in an intranasal mouse sepsis/lethality model. As the target indication for a preventive vaccine in humans is pharyngitis, an intranasal challenge model for the evaluation of candidate antigens is believed to be physiologically more relevant than an intravenous or intraperitoneal model, which have been described previously (Guzman et al., J. Inf. Dis. 179: 901 (1999); Stalhammar-Carlemalm et al., Mol. Microbiol. 33: 208 (1999)). Therefore protection was assessed in three distinct models, all applying the bacterial

challenge via the intranasal route. Protection was observed for 9 distinct proteins in the intranasal challenge model, some of which were tested as a fragment of the full length recombinant protein.

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Since protection against streptococcal challenge is mediated by antibodies, immunizations were first performed using CFA/IFA as adjuvant in order to obtain very high levels of antibodies. Subsequently, experiments were also performed with Alum and IC31TM as adjuvants, as these adjuvants are suited for use in humans and would be a preferred choice for a vaccine to prevent group A streptococcal infections in humans. As can be seen for the experiment depicted in Figure 1, fragment Spy0292-1 performed as well as full length Spy0292 protein for protection, while Spy0292-3 showed lower levels of protection. This clearly indicates that one region useful for protection lies within the sequence encompassing the Spy0292-1 protein.

Similar results were obtained for the proteins, Spy0269 (good protection also observed with Spy0269-1), Spy0416 (good protection also observed with Spy0416A-1, Spy0416A-6 and Spy0416A-7), and Spy0872 (good protection also observed with Spy0872-2).

For the proteins Spy0488, Spy0895, and Spy1727 full length recombinant proteins were used (Table 1), as these proteins have been shown for the first time to be capable to provide protection against lethal *S. pyogenes* challenge. Especially protein Spy0895 shows promise as a vaccine candidate, because it provided protection against group A streptococcal infection in all three models listed in Table 1.

Spy1536 and Spy1666 have been shown to provide protection in an intravenous challenge model before (WO 2004/078907), but importantly it could now be shown that they also provide protection in the physiologically more relevant intranasal challenge model. Spy1536 was most consistent in providing significant protection in all three models of GAS infection. Besides these two antigens, Spy0895 and Spy1536, several antigens showed protection in at least 2 models: Spy0269-1, Spy0292-1, Spy0416A-1, Spy0872-2, Spy1666 and Spy1727. Importantly, several antigens showed a level of protection that was as high as the level seen for the positive control protein M1 (e.g. Spy0416A-1, Spy0488, Spy0895; Table 1).

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These data clearly provide evidence, that the selected proteins are promising candidates for vaccine development. In addition, proteins Spy0269, Spy0292, Spy0416, and Spy0872 have been shown to possess amino acid sequences that are dispensable for protection, since sub-fragments were capable to provide the same or even superior levels of protection than the full length recombinant protein.

Table 1: Recombinant proteins of *S. pyogenes* and fragments thereof assessed for protection in murine models of infection.

ORFI Protein	Length ¹ (aa)	Amino acids ¹ (from – to)	SEQ ID No	Calculated MW (kDa) ²	Vector	Base pairs ¹ (from – to)	Protection ³
Spy0269	837	36 - 873	57	92.34	pET28b	106-2619	10% (30%, 60%) ^A
Spy0269-1	452	37-488	1	50.85	pET28b	109-1464	50% (10%, 50%)B,A,C
Spy0292	388	23 - 410	68	44.91	pET28b	67-1233	60% (10%, 90%) ^{A,C}
Spy0292-1	162	23-184	2	19.41	pET28b	67-554	56% (10%, 90%) ^{A,B}
Spy0292-3	278	23-300	3	32.39	pET28b	67-900	30% (10%, 90%) ^A
Spy0416A	834	34 - 867	89	95.80	pET28b	100-2601	20% (10%, 63%) ^A
Spy0416A-1	644	34-677	4	74.70	pET28b	100-2031	80% (20%, 80%) ^{C,A}
Spy0416A-6	311	148-458	5	38.77	pET28b	442-1374	40% (10%, 63%) ^A
Spy0416A-7	487	72-558	6	57.68	pET28b	214-1674	63% (10%, 63%) ^A
Spy0416B	882	736 - 1617	56	103.08	pET28b	2206-4851	20% (10%, 63%) ^A
Spy0488	331	1-331	8	37.84	pET28b	1-993	90% (20%, 80) ^{C,A}
Spy0872	613	28 - 640	120	68.38	pET28b	82-1920	20% (0%, 60%) ^A
Spy0872-2	290	351-640	7	33.02	pET28b	1051-1920	60% (0%, 60%)A,C,B
Spy0895	261	2-262	9	32.15	pET28b	4-786	90% (20%, 80%) ^{C,A,B}
Spy1536	314	32-345	131	35.27	pET28b	94-1035	70% (20%, 80%) ^{C,A,B}
Spy1666	315	23-337	132	37.02	pET28b	67-1011	60% (20%, 80%) ^{C,B}
Spy1727	263	1-263	10	32.43	pET28b	1-789	70% (20%, 80%) ^{C,B}

¹ Length, amino acids and base pairs are calculated for the S. pyogenes gene specific sequence only.

Brackets show protection in the respective model with the negative (PBS + adjuvant only) and positive control (M protein). If protection was seen in more than one model, the protection data of the model listed first are shown.

The calculated molecular weight includes amino acids derived from the vector and the His6-tag.

³ Protection is based on the animal model as indicated:

A s.c. immunization using CFA/IFA as adjuvant, i.n. challenge with S. pyogenes A20

B s.c. immunization using ALUM as adjuvant and i.n. challenge with S. pyogenes A20

C intranasal immunization using IC31[™] or a mucosal adjuvant and intranasal challenge with either S. pyogenes A20 or A147.

ORF-protein	Plasmid name	Primer ¹	Name	Restriction enzyme
SPy0269	pET28b-SPy0269	TAGTAGCCATGGGCGATGATAGAGCCTCA GGA SEQ ID NO: 21	210-2129	Ncol
		TAGTAGGCGGCCGCCTTAGATTCCTTACG GAACCT SEQ ID NO: 22	210-2196	Notl
SPy0269-1	pET28b-SPy0269-1	TAGTAGCCATGGGCGATGATAGAGCCTCA GGA SEQ ID NO: 23	210-2129	Ncol
		TAGTAGGCGGCCGCAACAGGCGCATTAGG G SEQ ID NO: 24	210-2719	Noti
SPy0292	pET28b-SPy0292	TAGTAGCCATGGGCGAAGAGTATTCGGTA ACTGC SEQ ID NO: 25	210-2131	Ncol
		TAGTAGGCGGCCGCTAAAGAGGTATTGAC ATACCT SEQ ID NO: 26	210-2197	Notl
SPy0292-1	pET28b-SPy0292-1	TAGTAGCCATGGGCGAAGAGTATTCGGTA ACTGC SEQ ID NO: 27	210-2131	Ncol
		TAGTAGGCGGCCGCGCAAAAACAATTTC ATCATC SEQ ID NO: 28	210-2954	Notl
SPy0292-3	pET28b-SPy0292-3	TAGTAGCCATGGGCGAAGAGTATTCGGTA ACTGC SEQ ID NO: 29	210-2131	Ncol
		TAGTAGGCGGCCGCTTCAATTAACTGGAC TTTTTG SEQ ID NO: 30	210-2956	Notl
SPy0416A	pET28b-SPy0416A	TAGTAGGAATTCGGCAGATGAGCA CAATG SEQ ID NO: 31	210-2246	EcoRI
		TAGTAGCTCGAGCTCTGAACCAAGAGTGA CAAG SEQ ID NO: 32 TAGTAGGAATTCGGCAGATGAGCTAAGCA	210-2247	Xhol
SPy0416A-1	pET28b-SPy0416A-1	CAATG SEQ ID NO: 33 TAGTAGCTCGAGTGCCCCTTGCTGACGCG	210-2246	EcoRi
		GTG SEQ ID NO: 34 TAGTAGGAATTCGGCAGTTATTGACACAGG	210-2663	Xhol
SPy0416A-6	pET28b-SPy0416A-6	G SEQ ID NO: 35 TAGTAGCTCGAGTAGGCTATCTTTTATGTC	210-2715	EcoRI
		SEQ ID NO: 36 TAGTAGGAATTCGTCACAAATCACTCTCAA	210-2717	Xhol
SPy0416A-7	pET28b-SPy0416A-7	G SEQ ID NO: 37 TAGTAGCTCGAGACTTCCTGTACCATTGCC	210-2716	EcoRI
		SEQ ID NO: 38 TAGTAGGAATTCGCATGTAGACCCACAAA	210-2718	Xhol
SPy0416B	pET28b-SPy0416B	AGGC SEQ ID NO: 39 TAGTAGCTCGAGCGTTGATGGTAGGCCTTT	210-2248	EcoRI
		TGC SEQ ID NO: 40 TAGTAGCCATGGGCTTGCGGCAGATTCAG	210-2249	Xhol
SPy0488	pET28b-SPy0488	TCCATT SEQ ID NO: 41 TAGTAGGCGGCCGCACTTTTTAACCTGTCC	210-2139	Ncol
		TCAGC SEQ ID NO: 42 TAGTAGCCATGGGCGATCAAGTTGATGTG	210-2199	Notl
SPy0872	pET28b-SPy0872	CAATTC SEQ ID NO: 43 TAGTAGGCGGCCGCTGTTATTGGAAGAGT	210-2143	Ncol
		GGAACT SEQ ID NO: 44 TAGTAGCCATGGGCGCTATAATAAATCATG	210-2144	Notl
SPy0872-2	pET28b-SPy0872-2	CT SEQ ID NO: 45 TAGTAGGCGGCCGCTGTTATTGGAAGAGT	210-2962	Ncol
SPy0895	pET28b-SPy0895	GGAACT SEQ ID NO: 46 TAGTAGCCATGGGCACTAATAATCAAACA	210-2144	Notl

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ORF-protein	Plasmid name	Primer 1	Name	Restriction enzyme
		CTA SEQ ID NO: 47		
		TAGTAGGCGGCCGCGACAATAGATTGTCT	210-2201	Notl
		CCAAAG SEQ ID NO: 48	210-2201	1400
00 4500	-FT205 CD-4526	TAGTAGCCATGGGCATTGAAATGCCTGGA	210-2161	Ncol
SPy1536	pET28b-SPy1536	GGCG SEQ ID NO: 49	210-2101	1,100,
		TAGTAGGCGGCCGCTTTGCGAAGATAAAC	210-2207	Notl
		CAGTGC SEQ ID NO: 50		
CD::4666	pET28b-SPy1666	TAGTAGCCATGGGCACAAAAGAATTTCATC	210-2165	Ncol
SPy1666	pe 1200-3Fy 1000	ACGTG SEQ ID NO: 51	2.02.00	
	17	TAGTAGGCGGCCGCTTTCCGAATTTTTTTG	210-2209	Notl
		GCAAC SEQ ID NO: 52		
CDv4727	pET28b-SPy1727	TAGTAGCCATGGGCGTGACAACGACGAA	210-2167	Ncol
SPy1727	p=1200-3Fy1727	CAAG SEQ ID NO: 53		
		TAGTAGGCGGCCGCTTTCTTTCTAAATATT	210-2210	Notl
		TCTCT SEQ ID NO: 54		

Primer, letters in bold indicate gene-specific sequences, letters underlined indicate the restriction enzyme sites, letters in normal font indicate sequences necessary for cloning, but not present in the final plasmid construct used for expression. The first primer always refers to the sense and the second primer to the anti-sense oligonucleotide in relation to the encoded gene used for amplification.

Example 2: Group A streptococcal antigens and variants thereof.

10 Experimental procedures

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Preparation of streptococcal genomic DNA

5 mL Todd-Hewitt Broth medium were inoculated with the respective strain of *S. pyogenes* (as listed in Table 3) from a frozen stab and grown without shaking at 37°C overnight. 4 mL of the culture were then harvested by centrifuging at 13,000 rpm in a biofuge fresco (Haereus) for 5 min and the supernatant was removed. DNA was isolated from the bacterial cell pellets following the protocol of Wizard® Genomic DNA Purification Kit (Promega). The DNA pellets were finally dried on air and dissolved in 70 μl ddH₂O.

PCR and sequence analyses of S. pyogenes genes

In order to determine the sequence of an antigen from diverse *S. pyogenes* strains, PCR was performed with primers specific for the gene of interest. *S. pyogenes* strains used for these analyses are shown in Table 3. Oligonucleotide sequences as primers for PCR were designed for the selected antigens in order to be able to amplify the full gene. Sequencing was performed with dedicated primers using the PCR products as templates. The sequences of the oligonucleotides are listed in Table 4. Genomic DNA of all *S. pyogenes*

strains was prepared as described above. PCR was performed in a reaction volume of 25 µl using Taq polymerase (1 U), 200 nM dNTPs, 10 pMol of each oligonucleotide and the kit according to the manufacturer's instructions (Invitrogen, The Netherlands). As standard, 30 cycles (1x: 5 min. 95°C, 30x: 30 sec. 95°C, 30 sec. 56°C, 120 sec. 72°C, 1x 4 min. 72°C) were performed, unless conditions had to be adapted for individual primer pairs. PCR samples were sequenced with the oligonucleotides as listed in Table 10. Sequencing was performed at Agowa (Germany).

Table 3: S. pyogenes clinical isolates utilized for the present study.

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No.	Strain	Country of origin	Serotype
1	Schmitz 1/94	Netherlands	1
2	Schmitz 1/12	Portugal	1
3	Schmitz 1/5	Portugal	11
4	Schmitz 2/14	Germany	1
5	Schmitz 1/74	England	3
6	Schmitz 1/35	Spain	3
7	Schmitz 1/41	France	3
8	RDN 78	unknown	3.1
9	Schmitz 1/17	Portugal	4
10	Schmitz 1/156	Switzerland	4
11	Schmitz 1/22	Spain	4
12	RDN 60	unknown	5
13	Schmitz 1/174	Austria	6
14	Schmitz 1/97	Belgium	6
15	Schmitz 1/29	Spain	9
16	Schmitz 1/92	Netherlands	11
17	Schmitz 1/39	Spain	12
18	Schmitz 1/248	Poland	12
19	Schmitz 1/59	England	12
20	RDN 02	unknown	19
21	Schmitz 1/76	England	22
22	Schmitz 1/177	Austria	22
23	Schmitz 1/43	France	22
24	Schmitz 2/32	Germany	22
25	RDN 136	unknown	22.2
26	Schmitz 1/136	Germany	25
27	Schmitz 1/56	France	28
28	Schmitz 1/108	Belgium	28
29	Schmitz 1/85	Netherlands	28
30	Schmitz 2/50	Germany	28
31	Schmitz 1/194	Italy	44
32	Schmitz 1/234	Turkey	44
33	Schmitz 1/103	Belgium	44
34	Schmitz 1/253	Poland	49
35	Schmitz 1/141	Germany	49
36	Schmitz 1/123	Germany	49

37	Schmitz 2/30	Germany	66 or 90
38	Schmitz 1/144	Germany	76
39	Schmitz 1/99	Belgium	78
40	RDN 120	unknown	81
41	Schmitz 1/142	Germany	83
42	Schmitz 1/176	Austria	83
43	Schmitz 1/25	Spain	83
44	RDN 75	unknown	85
45	Schmitz 2/46	Germany	89
46	Schmitz 2/9	Germany	90
47	Schmitz 2/23	Germany	90
48	RDN 116	unknown	94
49	Schmitz 1/55	France	118
50	Schmitz 1/68	England	118
51	Schmitz 1/3	Portugal	118

Table 4: Oligonucleotides used for sequence conservation analyses. Shown are the ORF and primer names, orientation of the primer relative to the gene, the sequence, and the position relative to the gene. Oligonucleotides were used for both PCR amplification of the gene or gene fragment and subsequent sequence analyses.

ORF	Primer name	Orientation	Sequence	SEQ ID NO:	Position relative to gene
	210-4752	sense	TGACCTTCAAATCATTGCTGA	209	-103 to -82
	210-4759	antisense	TTTTGCACTTCTGGTGTCAA	210	1014 to 1034
Spy0269	210-4754	sense	TTGCCAAAGCTAGTCCAGGT	211	931 to 951
-,,	210-4761	antisense	AGTATTATCAATGCGCTCACG	212	2028 to 2049
	210-4756	sense	AAAAGCTCATTTGCAATATCTAAGG	213	1967 to 1992
	210-4763	antisense	GCTGGTGAATCTGATTTTTCAA	214	2875 to 2897
	210-4575	sense	TCTTGTGAGGTAAGTCATTACCTTAG	215	-79 to -53
	210-4576	antisense	TTCATCATCTGGTTCTGTATTAGG	216	516 to 540
Spy0292	210-4577	sense	GGTCGTCAATTCAACTGGC	217	464 to 483
00,0202	210-4578	antisense	GCGATCATTGTGGATGATTTC	218	1031 to 1052
	210-4579	sense	AAACTGTCAAACTTGTAGCCC	219	946 to 967
	210-4580	antisense	TGTTAGGATTGGCCTAGTTTG	220	1304 to 1325
	210-4588	sense	TGAGTTAATGATTAACATTAAACTGGT	221	-56 to -29
	210-4591	antisense	TGACATAAGCAAATTGATGCG	222	1387 to 1408
	210-4592	sense	CCATCTATTCAGAGTCTGTCGAC	223	1327 to 1350
Spy0416	210-4595	antisense	CCTTGTCACTAGCATGGTAGAC	224	2802 to 2824
оруотто	210-4596	sense	TTGCAGCCTTCAAAGGTG	225	2749 to 2767
	210-4599	antisense	AAGACACATTACCAGCTCTATCTTC	226	4128 to 4153
	210-4600	sense	CAGATGGTTCTTACACCATTTC	227	4063 to 4085
	210-4603	antisense	AATCTCAAAGAAAGGTCAGACTG	228	4982 to 5005
	210-5497	sense	AAAGCTCGTCATTTTATATGATTT	229	-195 to -171
Spy0488	210-4767	antisense	TTTAATGAGAGTTGTCATTCGTTCA	230	497 to 522
Opyu-tou	210-4765	sense	TTTTCTTGTTCAACCGCAAG	231	404 to 424
	210-4766	antisense	GCGCTCACAGCTACTTCAGA	232	1052 to 1072
Spy0872	210-4581	sense	CAAAATCATAGTAAACTTGATCTATAAG	CG 233	-55 to -26
	210-4584	antisense	GAAGAATTAGTTGCAGTTCCG	234	1103 to 1124
	210-4585	sense	GTTGCTGTAGCACCAGGTATC	235	1005 to 1026

	210-4587	antisense	CCAGCACGAATTAGATCATCTAG	236	2111 to 2134
Spy0985	210-4768	sense	CTGAAGAGCGCCAAACAACT	237	-63 to -43
	210-4771	antisense	TCGAAGAAGTAACCTTTGATTAATGT	238	864 to 890
Spy1536	210-4772	sense	GCTCTAGTCGTGTGAGAGAGCTAA	239	-90 to -66
Эру 1550	210-4775	antisense	TGTCTATCTGGTTCAACCGTTTT	240	1089 to 1112
Spy1666	210-4780	sense	GTGGCTAAGTCAGTGCTTGCT	241	-80 to -59
Ору 1000	210-4783	antisense	AAGTTTTTATTCGTTTTTGCAAGG	242	1055 to 1079
Spy1727	210-4776	sense	GATCATTGACTAAGTAGCCTAAAACAA	243	-76 to -49
Opy 1727	210-4779	antisense	CCAAAAACGTCATGCCAAC	244	879 to 898

RESULTS

Gene conservation analysis of selected streptococcal antigens

The PCR and sequencing of the 9 selected genes was performed as described under Methods. Table 3 shows the strains used for sequencing, while Table 4 lists the oligonucleotides employed for the PCR and sequencing analyses.

Sequence analyses of Spy0269

- Sequences were obtained from all 51 strains. The level of amino acid sequence identity ranged from 98.7% to 100% as compared to the sequence of Spy0269 from *S. pyogenes* SF370. Table 5 lists all 36 amino acid positions which showed a distinct amino acid as compared to Spy0269 from *S. pyogenes* SF370.
- Table 5: Gene conservation of Spy0269. 1, observed amino acid at respective position in any of the sequenced genes of the respective S. pyogenes strains.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	AA change ²	Strains with the respective change ¹	Strains with the respective change ²
30	30	V			Schm1_142, Schm1_177, Schm1_43, RDN75	
68	68	D	E		Schm1_76, Schm1_92, Schm1_142, Schm1_176, Schm1_177, Schm1_25, Schm1_43, Schm2_32, RDN136, RDN75	
73	73	Т	A		Schm1_142, Schm1_177, Schm1_43	

80 80 E K Schml 55, Schml 68, Schml 3, Schml 23, Schml 23, Schml 23, Schml 50,					
Schml	80	80	F	K	Schm1 55.
Schml 3, Schml 23, Schml 23, Schml 23, Schml 23, Schml 23, Schml 30, Schml 17, Schml 159, Schml 197, Schml 197, Schml 197, Schml 141, Schml 141, Schml 143, Schml 144, Schml 144, Schml 144, Schml 144, Schml 144, Schml 150, Schml 246, Schml 246, Schml 250, Schml 2			_	'`	
Schm2 23 Schm2 30				}	
Schm2 30					
83 83 E K Schml 59 Schml 59 Schml 59 Schml 59 Schml 59 Schml 142 Schml 142 Schml 142 Schml 143 Schml 143 Schml 143 Schml 144 Schml 24 Schml 25 Schml 25 Schml 26 Schml 27 Schml 27 Schml 27 Schml 28 Schml 29 Schml 20 Schml 27 Schml 27 Schml 27 Schml 28 Schml 27 Schm					_
Schm1 97 Schm1 97 Schm1 97 Schm1 97 Schm1 97 Schm1 142 Schm1 147 Schm1 147 Schm1 99 Schm2 14 Schm2 46 Schm2 46 Schm1 35 Schm1	83	83	F	K	
Schm1 142, Schm1 142, Schm1 142, Schm1 143, Schm1 143, Schm1 143, Schm1 143, Schm2 14, Schm2 14, Schm2 14, Schm2 14, Schm1 141, Schm1 141, Schm1 141, Schm1 141, Schm1 141, Schm1 141, Schm2 9, Schm2 9, Schm2 9, Schm2 9, Schm2 9, Schm2 9, Schm1 103, Schm1 104, Schm1 17, Schm1 17, Schm1 17, Schm1 142, Schm1 142, Schm1 142, Schm1 142, Schm1 143, Schm1 108, Schm1 177, Schm1 108, Schm1 177, Schm1 103, Schm1 104, Schm2 20, Schm1	00		_	'`	
94 94 E K Schm1_142, Schm1_177, Schm1_99, Schm1_99, Schm2_46 150 150 A V Schm1_35, Schm1_141, Schm2_46 150 230 230 A G Schm1_141, Schm2_9, Schm2_9, Schm2_99, Schm1_35, Schm2_99, Schm1_156, Schm1_156, Schm1_156, Schm1_156, Schm1_159, Schm1_168, Schm1_168, Schm1_179, Schm2_30, Schm1_179, Schm1_199, Schm1_199, Schm1_130, S))	
Schm1_177, Schm1_43 97	94	04		- k	
Schm1 43 Schm1 99 Schm2 14 Schm2 14 Schm2 46 Schm1 35 Schm1 141 Schm2 99 Schm2 14 Schm1 35 Schm1 141 Schm1 155 Schm1 103 Schm1 103 Schm1 103 Schm1 108 Schm1 109 Schm1 141 Schm1 176 Schm1 176 Schm1 177 Schm1 192 Schm1 141 Schm1 176 Schm1 177 Schm1 178 Schm1 178 Schm1 179 Schm1 179 Schm1 179 Schm1 170 S	34	34	<u> </u>		
97 97 H N Schm1 99, Schm2 14, Schm2 14, Schm2 46 150 150 A V Schm1 74, Schm1 35, Schm1 174, Schm1 174, Schm1 174, Schm1 174, Schm1 174, Schm1 35 230 230 A G Schm1 35 249 249 E D Schm1 103 276 276 A V Schm1 108 279 279 G D Schm1 108 279 279 G D Schm1 108 3 Schm2 23, Schm2 30 307 307 A G Schm1 92 3 Schm2 30 307 107 A G Schm1 176, Schm1 186, Schm1 176, Schm1 192, Schm1 192, Schm1 177, Schm1 192, Schm1 177, Schm1 177, Schm1 177, Schm1 177, Schm1 178,					
Schm2_14, Schm2_46	07	07			
Schm2_46	97	97		IN I	
150				1	
Schm1_35, Schm1_141, Schm1_174, Schm1_174, Schm1_174, Schm1_41, Schm2_50, RDN60, RDN75, Schm2_50, RDN60, RDN75, Schm1_35, Schm1_35, Schm1_103, Schm1_103, Schm1_108, Schm1_108, Schm1_108, Schm1_108, Schm1_108, Schm1_108, Schm1_23, Schm2_30, Schm1_17, Schm1_176, Schm1_176, Schm1_176, Schm1_174, Schm1_174, Schm1_174, Schm1_174, Schm1_177, Schm1_176, Schm1_177, Schm1_178, Schm1_178, Schm1_178, Schm1_179, Schm1_199, Schm1	450	450			
Schm1_141, Schm1_174, Schm1_141, Schm2_9, Schm2_50, RDN60, RDN78, RDN75, Schm1_166, Schm1_166, Schm1_156, Schm1	150	150	A	V	
Schm1 174, Schm2 9, Schm2 50, RDN60, RDN78, RDN75			1		
Schm1_41, Schm2_9, Schm2_9, Schm2_9, Schm2_9, Schm2_9, Schm2_50, RDN60, RDN78, RDN75		-		1	
Schm2_90, Schm2_50, RDN60, RDN78, RDN78, RDN75, RDN76, RDN75, R					
Schm2_50, RDN60, RDN78, RDN60, RDN78, RDN60, RDN78, RDN75 230 230]				
RDN60, RDN78, RDN75					
RDN75				-	
230 230			•		
249					
276	230	230	A	G	Schm1_35
276	249	249	E	D	Schm1_103
Schm1 108 Schm1 108 Schm1 108 Schm1 55 Schm1 56 Schm1 58 Schm1 30 Schm2 20 Schm2 30 Schm1 17 Schm1 17 Schm1 17 Schm1 17 Schm1 176 Schm1 125 Schm1 141 Schm1 174 Schm1 174 Schm1 177 Schm1 17 Schm1 17 Schm1 19 Schm1 19 Schm1 19 Schm1 19 Schm1 19 Schm1 19 Schm1 136 Schm2 14 Schm2 32 Schm2 46 Schm2 46 Schm2 50 RDN60 RDN02 RDN136 RDN120 RDN116 RDN116 Schm1 155 Schm1 155 Schm1 156 Schm1 128 Schm1 128 Schm1 128 Schm1 156 Schm1 156 Schm1 156 Schm1 156 Schm1 156 Schm1 156 Schm1 128 Schm1 156 Schm1 156 Schm1 128 Schm1 156 Schm1 156 Schm1 156 Schm1 156 Schm1 156 Schm1 128 Schm1 156 Schm1 156 Schm1 156 Schm1 158 Schm1 156 Schm1 158 Schm1	276	276	A		
279 C					
Schm1	279	279	G	D	
Schm1_3, Schm2_23, Schm2_30 307	2.0	2.0			
Schm2 23, Schm2 30 Schm1 92]	
Schm2 30 Schm1 92	1	İ	l		
307 307 A G Schm1 92 482 H R Schm1 177, 482 Schm1 56, 5chm1 76, 5chm1 92, 5chm1 182, 5chm1 183, 5chm1 184, 5chm1 174, 5chm1 176, 5chm1 177, 5chm1 25, 5chm1 25, 5chm1 99, 5chm1 99, 5chm1 199, 5chm1 123, 5chm1 136, 5chm1 136, 5chm2 14, 5chm2 32, 5chm2 46, 5chm2 50, RDN60, RDN02, RDN136, RDN120, RDN116 485 485 N K Schm1 39, 5chm1 39, 5chm1 39, 5chm1 39, 5chm1 123, 5chm1 186, 5chm1 248,					
## ## ## ## ## ## ## ## ## ## ## ## ##					Sobm2 20
Schm1_56, Schm1_76, Schm1_76, Schm1_92, Schm1_142, Schm1_123, Schm1_141, Schm1_177, Schm1_177, Schm1_25, Schm1_25, Schm1_18, Schm1_177, Schm1_99, Schm1_123, Schm1_136, Schm2_14, Schm2_14, Schm2_14, Schm2_46, Schm2_46, Schm2_50, RDN60, RDN60, RDN16, RDN16, RDN116, RDN116, RDN116, RDN116, Schm1_55, Schm1_55, Schm1_68, Schm1_55, Schm1_68, Schm1_56, Schm1_5248,	207	207			
Schm1_76, Schm1_92, Schm1_142, Schm1_253, Schm1_08, Schm1_141, Schm1_174, Schm1_176, Schm1_177, Schm1_25, Schm1_25, Schm1_97, Schm1_97, Schm1_99, Schm1_123, Schm1_136, Schm1_136, Schm2_14, Schm2_14, Schm2_32, Schm2_46, Schm2_50, RDN160, RDN02, RDN166, RDN120, RDN166, RDN120, RDN116 Schm1_39, Schm1_39, Schm1_55, Schm1_68, Schm1_68, Schm1_156, Schm1_68, Schm1_156, Schm1_1248,					Schm1_92
Schm1_92, Schm1_142, Schm1_143, Schm1_108, Schm1_174, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_25, Schm1_25, Schm1_97, Schm1_99, Schm1_97, Schm1_99, Schm1_136, Schm1_136, Schm2_14, Schm2_32, Schm2_14, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN120, RDN136, RDN120, RDN116					Schm1_92 Schm1_17,
Schm1_142, Schm1_253, Schm1_108, Schm1_141, Schm1_174, Schm1_174, Schm1_176, Schm1_177, Schm1_25, Schm1_43, Schm1_99, Schm1_99, Schm1_99, Schm1_123, Schm1_136, Schm2_14, Schm2_14, Schm2_32, Schm2_46, Schm2_46, Schm2_50, RDN60, RDN00, RDN136, RDN136, RDN120, RDN116					Schm1_92 Schm1_17, Schm1_56,
Schm1_253, Schm1_108, Schm1_114, Schm1_174, Schm1_174, Schm1_176, Schm1_176, Schm1_25, Schm1_25, Schm1_99, Schm1_99, Schm1_99, Schm1_136, Schm1_136, Schm2_14, Schm2_32, Schm2_46, Schm2_14, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN00, RDN16, RDN120, RDN116 485					Schm1_92 Schm1_17, Schm1_56, Schm1_76,
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		R	K	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9,
		R	K	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32,
		R	K	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136,
		R	K	Schm1_174 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78
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636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_36, RDN78 Schm1_74, Schm1_74, Schm1_74, Schm1_76,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_75, Schm1_74, Schm1_75, Schm1_75, Schm1_75, Schm1_75, Schm1_35,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_36, RDN78 Schm1_74, Schm1_74, Schm1_74, Schm1_76,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_141, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_176,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_176, Schm1_176, Schm1_176, Schm1_176, Schm1_176, Schm1_176, Schm1_176, Schm1_25,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_76, Schm1_76, Schm1_76, Schm1_176, Schm1
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm2_32, RDN136, RDN78 Schm1_74, Schm2_9, Schm2_35, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_76, Schm1_176, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_74, Schm1_76, Schm1_76, Schm1_176, Schm1_25, Schm1_176, Schm1_25, Schm1_176, Schm1_25, Schm1_29, Schm2_32,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_76, Schm1_76, Schm1_176, Schm1_25, Schm1_176, Schm1_25, Schm1_176, Schm1_27, Schm2_32, RDN136,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_74, Schm1_76, Schm1_76, Schm1_176, Schm1_25, Schm1_176, Schm1_25, Schm1_176, Schm1_25, Schm1_29, Schm2_32,
636	636	A A	M	Schm1_174 Schm1_74, Schm1_76, Schm1_35, Schm1_176, Schm1_25, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_76, Schm1_176, Schm1_176, Schm1_176, Schm1_25, Schm1_41, Schm2_9, Schm2_32, RDN136, RDN78 Schm1_74, Schm1_74, Schm1_76, Schm1_76, Schm1_176, Schm1_25, Schm1_176, Schm1_25, Schm1_176, Schm1_27, Schm2_32, RDN136,

				Schm1_108
650	650	V	E	Schm2_9
666	666	F	L	Schm1_22
700	700	Α	T	Schm1_17,
	· ·			Schm1_39,
				Schm1_55,
				Schm1_56,
				Schm1_253,
i		•		Schm1_68,
1				Schm1_108,
				Schm1_156,
			1	Schm1_248,
				Schm1_3,
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				Schm1_29,
i				Schm1_59,
}				Schm1_97,
				Schm1_123,
				Schm1_136,
]				
}				Schm2_23,
Ţ			[Schm2_30,
				RDN02,
				RDN120,
				RDN116
703	703	Α	V	Schm2_50,
703	703	^		
710	7/0			RDN60
710	710	S	G	Schm1_17,
				Schm1_59,
V				Schm1_97
733	733	E	G	Schm1_56,
100	, 00	_		Schm1_108
750	750			
750	750	Α	P	Schm1_22
752	752	P	S	Schm1_55,
				Schm1_74,
				Schm1_76,
1			1	Schm1_92,
f				Schm1_142,
				Schm1_144,
}				Schm1_194,
				Schm1_35,
				Schm1_68,
				Schm1_176,
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				Schm1_177,
				Schm1_234,
1				Schm1_3,
ſ				Schm1_25,
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				Schm1_41,
				Schm1_41, Schm1_43,
				Schm1_41, Schm1_43, Schm1_99,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_30, Schm2_32,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136,
				Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46,
758	758	P		Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN78
758 764	758 764	P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN78
758 764	758 764	P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN78 Schm1_92 Schm1_74,
		P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN136, RDN78 Schm1_92 Schm1_74, Schm1_76,
		P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN136, RDN78 Schm1_92 Schm1_74, Schm1_76, Schm1_92,
		P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN136, RDN78 Schm1_92 Schm1_74, Schm1_76,
		P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN136, RDN78 Schm1_92 Schm1_74, Schm1_76, Schm1_92, Schm1_142,
		P	L	Schm1_41, Schm1_43, Schm1_99, Schm1_103, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN136, RDN136, RDN78 Schm1_92 Schm1_74, Schm1_76, Schm1_92,

					Schm1_35,	
}	1		1	:	Schm1_176,	1
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	i			•	Schm1_234,	
	Į į			Į	Schm1_25,	
				j		
					Schm1_41,	
					Schm1_43,	ļ
				1	Schm1_99,	
					Schm1_103,	
Ì	ĺ		ì	Ì	Schm2_9,	1
			1		Schm2_14,	
			Į.	Į.	Schm2_32,	
				1	Schm2_46,	
1			1	1	RDN136,	}
1						
705	705		 	 	RDN78	
765	765	D	E		Schm1_74,	
					Schm1_76,)
					Schm1_92,	
			1	1	Schm1_142,	
					Schm1_144,	
			1		Schm1_194,	
			1	•	Schm1_35,	
					Schm1_176,	
1				1		
				1	Schm1_177,	
}		1	1	1	Schm1_234,	1
			}	}	Schm1_25,	1
					Schm1_41,	
					Schm1_43,	
)				}	Schm1_99,	}
					Schm1_103,	
Į i					Schm2_9,	
	•				Schm2_14,	
1	{		{	{	Schm2_32,	1
1					Schm2_46,	
ľ					RDN136,	1
		· 			RDN78	
794	794	L	F	H	Schm1_22	Schm2_23,
		 _				Schm2_30
873	873	K	R		Schm1_55,	
]			Į.	į.	Schm1_74,	
]			1		Schm1_76,	{
[Schm1_92,	
`	1			1	Schm1_142,	1
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!				1	Schm1_194,	
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]					Schm1_35,	
,					Schm1_68,	
]				J	Schm1_141,	
					Schm1_174,	
					Schm1_176,	
					Schm1_177,	
, l				l	Schm1_234,	[
j i				1	Schm1_3,	1
					Schm1_25,	
1	1				Schm1_23,	
1			1	1	Schm1_43,	}
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					Schm1_103,	
Į Į	ļ				Schm2_9,	ļ
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,			1	1	Schm2_23,	1
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			1		Schm2_30,	
					Schm2_30,	
					Schm2_30, Schm2_32,	
					Schm2_30,	

	RDN60,
	RDN136,
1	RDN78, RDN75

Sequence analyses of Spy0292

Sequences were obtained from all 51 strains. The level of amino acid sequence identity ranged from 97.3% to 100% as compared to the sequence of Spy0292 from *S. pyogenes* SF370. Table 6 lists all 36 amino acid positions which showed a distinct amino acid as compared to Spy0292 from *S. pyogenes* SF370.

Table 6: Gene conservation of Spy0292. ¹, observed amino acid at respective position in any of the sequenced genes of the respective S. pyogenes strains. ², second possible amino acid observed at the respective position. ³, third possible amino acid observed at the respective position.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	AA change²	AA change³	Strains with respective change ¹	Strains with respective change ²	Strains with respective change ³
21	21	S	N			Schm1 136		
32	32	Α	V			RDN02		
45	45	E	K			RDN60		
48	48	Α	Т			Schm1_56, Schm1_108, Schm1_85		
50	50	E	К			RDN75		
57	57	V	1			Schm2_50		
58	58	S	T			Schm2_50		
65	65	L	М			Schm1_141, Schm1_156, Schm1_174		
68	68	K	Q	N		Schm2 30	Schm2_50	
88	88	Y	D			Schm2_30		
89	89	E	D			Schm2_30		
93	93	N	Y			Schm2_50		
95	95	Т	S			Schm2_30		
96	96	1	M			Schm2_30		
101	101	L	Р			Schm2_30	;	
121	121	N	1			Schm2_50		
122	122	S	T			Schm2_50		
128	128	Α	Р	S		RDN60	RDN60	
137	137	K	N			Schm2_30		
141	141	K	E	Q		Schm1_17	Schm2_50	
147	147	R	L	W		Schm1_17	Schm2_50	RDN60
148	148	Q	L			Schm2_30, RDN60		
152	152	S	F			RDN120		
154	154	A	Т			Schm1_55, Schm1_68, Schm1_3, Schm1_29, Schm2_23, Schm2_30		
165	165	H	 	 		RDN60		

199 l	188 J	1 1	F	1	Schm1_174	1	
188 189	189	A	P		Schm1_174		
190	190		V		Schm1_253,		
	,				Schm1_123		
214	214	Α	D		Schm1_39,		
		i			Schm1_55,		
					Schm1_56,		
					Schm1_76,		
Į.	ļ			ļ .	Schm1_92,	ļ	
					Schm1_142, Schm1_144,		
		l			Schm1_108,		
		ľ]	Schm1_141,		
İ					Schm1_156,		
					Schm1_174,		
					Schm1_176,		
		i i			Schm1_177,		
					Schm1_234,		
		1			Schm1_248,		
					Schm1_25,		
1					Schm1_43,		
					Schm1_59,		
					Schm1_85, Schm1_99,		
					Schm1_103,		
					Schm2_32,		
)					Schm2_46,		
				f l	Schm2_50,		
		1			RDN60,		
					RDN02,		
}				}	RDN136,		
					RDN120		
240	240	V	l		Schm1_92,		
266	266	1			RDN120 Schm1_144,		
266	266	L			Schm1_144,		
					Schm1_103		
309	309	Y	S		Schm1_17,		
					Schm1_39,		
Į.		[Schm1_55,		
		i]			Schm1_56,		
		, ,	i		Schm1_74,		
}				1			
					Schm1_76,		
					Schm1_76, Schm1_92,		
					Schm1_76, Schm1_92, Schm1_142,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_141, Schm1_174, Schm1_174, Schm1_176,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_176,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_234,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_234, Schm1_248,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_141, Schm1_176, Schm1_176, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_248, Schm1_22,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_108, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_234, Schm1_248, Schm1_248, Schm1_25,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_108, Schm1_108, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_234, Schm1_234, Schm1_248, Schm1_248, Schm1_248, Schm1_25, Schm1_29,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_108, Schm1_108, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_234, Schm1_234, Schm1_248, Schm1_248, Schm1_248, Schm1_25, Schm1_29, Schm1_41,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_108, Schm1_108, Schm1_141, Schm1_174, Schm1_176, Schm1_177, Schm1_177, Schm1_234, Schm1_248, Schm1_248, Schm1_248, Schm1_248, Schm1_25, Schm1_25, Schm1_29, Schm1_41, Schm1_43,		
					Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_35, Schm1_108, Schm1_108, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_234, Schm1_234, Schm1_248, Schm1_248, Schm1_248, Schm1_25, Schm1_29, Schm1_41,		

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				Schm1_99, Schm1_103, Schm1_136, Schm2_9, Schm2_23, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN136, RDN120, RDN120, RDN75, RDN116
314	314	Р	S	Schm1_17, Schm1_22, Schm1_97
351	351	Α	Р	Schm1_177
371	371	G	Α	Schm1_234
386	386	Q	Н	Schm1_234

Sequence analyses of Spy0416

Sequences were obtained from all 50 strains excluding strain Schmitz 1/74. The level of amino acid sequence identity ranged from 98.1% to 100% as compared to the sequence of Spy0416 from *S. pyogenes* SF370. Table 7 lists all 103 amino acid positions which showed a distinct amino acid as compared to Spy0416 from *S. pyogenes* SF370. The gene showed in addition an insertion of 2 amino acids after position 31, as well as several deletions of amino acids at the indicated positions (e.g. strains Schmitz 1/17 and Schmitz 1/39).

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Table 7: Gene conservation of Spy0416. ¹, observed amino acid at respective position in any of the sequenced genes of the respective *S. pyogenes* strains. ², second possible amino acid observed at the respective position. Deletion or insertion refers to a missing or additional amino acid relative to Spy0416 of *S. pyogenes* SF370.

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Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	AA change ²	Strains with respective change ¹	Strains with respective change ²
21	21	1	V		Schm1_99, Schm2_46	
27	27	V	M		Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_142, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_141,	

1	1	1	i	Schm1_174,
				Schm1_176,
1	}			Schm1_177,
				Schm1_248,
	ļ			Schm1_3,
İ				Schm1_22,
				Schm1_25,
		ì	1	Schm1_29,
				Schm1_41,
1		1		Schm1_43,
				Schm1_59,
				Schm1_85,
				Schm1_97,
i				Schm1_99,
1		1		Schm1_123,
				Schm1_136,
	,	l l		Schm2_9,
				Schm2_14,
				Schm2_23,
				Schm2_30,
				Schm2_32,
1				Schm2_46,
				Schm2_50,
				RDN60,
				RDN136,
				RDN78,
				RDN120,
				RDN75,
				RDN116
29	29	Т	М	Schm1_17,
20	20	·		Schm1_39,
				Schm1_76,
				Schm1_142,
. ,				
				Schm1_35,
				Schm1_35, Schm1_141,
				Schm1_35, Schm1_141, Schm1_156,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136,
				Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75
Insertion	32	_	T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_136, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39,
Insertion	32			Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_76,
Insertion	32			Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_17, Schm1_39, Schm1_76, Schm1_142,
Insertion	32			Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_136, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_39, Schm1_76, Schm1_142, Schm1_35,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_39, Schm1_142, Schm1_142, Schm1_141,
Insertion	32			Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_17, Schm1_142, Schm1_141, Schm1_156,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_17, Schm1_39, Schm1_142, Schm1_141, Schm1_141, Schm1_156, Schm1_141, Schm1_156, Schm1_174,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_17, Schm1_35, Schm1_141, Schm1_141, Schm1_156, Schm1_174, Schm1_174, Schm1_174,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_76, Schm1_141, Schm1_141, Schm1_156, Schm1_174, Schm1_174, Schm1_174, Schm1_176, Schm1_176, Schm1_176, Schm1_177,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_176, Schm1_141, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_174, Schm1_176, Schm1_176, Schm1_176, Schm1_177, Schm1_176, Schm1_177, Schm1_177, Schm1_177, Schm1_248,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_176, Schm1_141, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_176, Schm1_174, Schm1_176, Schm1_176, Schm1_176, Schm1_176, Schm1_177, Schm1_176, Schm1_177, Schm1_177, Schm1_248, Schm1_248, Schm1_224,
Insertion	32		T	Schm1_35, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78, RDN75 Schm1_17, Schm1_39, Schm1_176, Schm1_141, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_174, Schm1_176, Schm1_176, Schm1_176, Schm1_177, Schm1_176, Schm1_177, Schm1_177, Schm1_177, Schm1_248,

Insertion	33	-	Schm1_43, Schm1_59, Schm1_97, Schm1_136, Schm2_9, Schm2_14, RDN136, RDN78 Schm1_17, Schm1_17, Schm1_22, Schm1_97
38	40	S	Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_142, Schm1_253, Schm1_35, Schm1_35, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_22, Schm1_22, Schm1_25, Schm1_59, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_97, Schm1_99, Schm1_103, Schm1_103, Schm1_136, Schm2_9, Schm2_14, Schm2_9, Schm2_14, Schm2_9, Schm2_14, Schm2_30, Schm2_30, Schm2_30, Schm2_31, Schm2_30, Schm2_31, Schm2_30, Schm2_31, Schm2_50, RDN136, RDN78, RDN16
40	42	M	Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_142, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_108, Schm1_174, Schm1_174, Schm1_176, Schm1_177,

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				Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_30, Schm2_30, Schm2_32, Schm2_32, Schm2_32, Schm2_36, RDN136, RDN136, RDN116	
49	51	A		Schm1_39, Schm1_76, Schm1_142, Schm1_35, Schm1_176, Schm1_177, Schm1_248, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm2_9, Schm2_14, Schm2_14, Schm2_32, RDN60, RDN136, RDN78	
54	56	Q	P	Schm1_55, Schm1_68, Schm1_3, Schm1_29, Schm2_23, Schm2_30	
55	57	Н	P	Schm1_55, Schm1_253, Schm1_68, Schm1_3, Schm1_29, Schm1_99, Schm1_123, Schm2_23, Schm2_30, Schm2_32, Schm2_46, RDN116	
67	69	K	Q	Schm1_17, Schm1_55, Schm1_56, Schm1_253, Schm1_68, Schm1_108,	

				Schm1_3, Schm1_22, Schm1_29, Schm1_97, Schm1_99, Schm1_123, Schm1_136, Schm2_23, Schm2_30, Schm2_30, Schm2_32, Schm2_32, Schm2_46, Schm2_50, RDN116	
68	70	S	P	Schm1_39, Schm1_55, Schm1_76, Schm1_142, Schm1_35, Schm1_68, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_177, Schm1_248, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_43, Schm1_43, Schm1_43, Schm1_59, Schm1_59, Schm2_9, Schm2_14, Schm2_15, Schm2_16,	Schm1_92
69	71	Q	P	Schm1_17, Schm1_56, Schm1_253, Schm1_108, Schm1_22, Schm1_85, Schm1_97, Schm1_99, Schm1_123, Schm1_136, Schm1_136, Schm2_32, Schm2_32, Schm2_46, Schm2_50, RDN116	
71	73	Т	I	Schm1_253, Schm1_123, Schm2_32	
74	76		V	Schm1_55, Schm1_253, Schm1_68, Schm1_3, Schm1_29, Schm1_99, Schm1_123, Schm1_136,	

		1	1	Cohm2 22
				Schm2_23, Schm2_30,
				Schm2_46
76	78	L	Р	Schm1_17,
				Schm1_55,
				Schm1_56,
				Schm1_92,
				Schm1_144, Schm1_194,
1				Schm1_154, Schm1_253,
				Schm1_68,
				Schm1_108,
				Schm1_141,
				Schm1_156,
				Schm1_174,
		l l	į	Schm1_234,
				Schm1_3, Schm1_22,
				Schm1_22, Schm1_29,
				Schm1_85,
				Schm1_97,
				Schm1_99,
				Schm1_123,
				Schm1_136,
				Schm2_23,
				Schm2_30, Schm2_46,
				Schm2_50,
				RDN60,
				RDN02,
				RDN116
77	79	K	Е	Schm1_55,
				Schm1_253,
				Schm1_68, Schm1_3,
				Schm1_3, Schm1_29,
				Schm1_99,
		1	•	Schm1_123,
				Schm1_136,
			1	Schm2_23,
				Schm2_30,
				Schm2_46
78	80	Т	1	Schm1_56, Schm1_108,
				Schm1_108, Schm1_85,
				Schm2_50
85	87	S	Р	Schm1_17,
]			Schm1_39,
				Schm1_55,
	· ·			Schm1_56,
				Schm1_76, Schm1_92,
1				Schm1_92, Schm1_142,
				Schm1_253,
				Schm1_35,
				Schm1_68,
		1		Schm1_108,
				Schm1_176,
		1	1	Schm1_177,
		1		Schm1_248, Schm1_3,
	1			Schm1_3, Schm1_22,
			1	Schm1_25,
				Schm1_29,
				Schm1_41,

				Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_14, Schm2_23, Schm2_23, Schm2_30, Schm2_50, RDN60, RDN136, RDN78
87	89		G	Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_92, Schm1_142, Schm1_1253, Schm1_35, Schm1_108, Schm1_176, Schm1_177, Schm1_177, Schm1_248, Schm1_3, Schm1_2, Schm1_25, Schm1_29, Schm1_41, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_97, Schm1_103, Schm1_103, Schm1_136, Schm1_136, Schm2_9, Schm2_14, Schm2_9, Schm2_30, Schm2_30, Schm2_50, RDN60, RDN136, RDN78
91	93	E	К	Schm1_99, Schm2_46,
93	95	Т	Deletion	RDN116 RDN60
102	104	A	S	RDN120, RDN75, RDN116
104	106	S	P	Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_142, Schm1_253, Schm1_35, Schm1_68, Schm1_108,

107 109 N Deletion Schml 92 110 112 S P Schml 17, Schml 39, Schml 56, Schml 76, Schml 76, Schml 76, Schml 92, Schml 142, Schml 142, Schml 108, Schml 1141, Schml 156, Schml 174, Schml 177, Schml 177, Schml 248, Schml 22, Schml 25, Schml 41, Schml 25, Schml 41, Schml 39, Schml 85, Schml 97, Schml 99, Schml 85, Schml 190, Schml 103, Schml 103, Schml 103, Schml 1136, Schml 123, Schml 136, Schml 124, Schml 136, Schml 1					Schm1_176, Schm1_177, Schm1_248, Schm1_3, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_99, Schm1_123, Schm1_136, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN136, RDN78, RDN120, RDN75, RDN116
	110	109	S	P	Schm1_39, Schm1_56, Schm1_76, Schm1_92, Schm1_142, Schm1_253, Schm1_35, Schm1_108, Schm1_141, Schm1_176, Schm1_176, Schm1_177, Schm1_248, Schm1_22, Schm1_25, Schm1_41, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_97, Schm1_103, Schm1_103, Schm1_136, Schm2_9, Schm2_14, Schm2_9, Schm2_14, Schm2_32, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN136, RDN78, RDN78, RDN78, RDN78, RDN78,

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215	217	E	G		Schm1_17, Schm1_92, Schm1_22, Schm1_97, Schm1_99, Schm2_46, RDN116	
228	230	A	Deletion		Schm1_17, Schm1_56, Schm1_92, Schm1_108, Schm1_22, Schm1_85, Schm1_97, Schm2_50, RDN120	
229	231	E	Deletion	D	Schm1_17, Schm1_56, Schm1_92, Schm1_108, Schm1_22, Schm1_85, Schm1_97, Schm2_50, RDN120, RDN116	Schm1_144, Schm1_194, Schm1_253, Schm1_234, Schm1_99, Schm1_123, Schm1_136, Schm2_46, RDN02
230	232	Α	Deletion		RDN116	
238	240	Н	N		Schm1_17, Schm1_92, Schm1_22, Schm1_97	
273	275	D	E		Schm1_92, Schm1_99, Schm2_46, RDN120, RDN116	
308	310	Α	Т		Schm1_56, Schm1_108, Schm1_85, Schm2_50	
320	322				Schm1_17, Schm1_39, Schm1_56, Schm1_56, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_176, Schm1_234, Schm1_248, Schm1_248, Schm1_248, Schm1_248, Schm1_248, Schm1_248, Schm1_25, Schm1_25, Schm1_29, Schm1_41,	

				Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
428	430	Т	Α	Schm1_142
429	431	V	A	Schm1_17, Schm1_22, Schm1_97
431	433	E	G	Schm1_253, Schm1_123
434	436	N	S	RDN116
449	451	V	F	Schm1_177
453	455	S	N	Schm1_142, Schm1_35, Schm1_141, Schm1_174, Schm1_176, Schm1_177, Schm1_248, Schm1_25, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_97, Schm1_123, Schm1_136, Schm1_136, Schm2_9, RDN136 Schm1_177,
463	465			RDN136
478	480	N	K	Schm1_17, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_176, Schm1_177, Schm1_234, Schm1_23, Schm1_25, Schm1_43, Schm1_97, RDN60, RDN02, RDN136, RDN120, RDN116

481	483	D	N	Schm1_55, Schm1_68, Schm1_3, Schm1_29, Schm1_136, Schm2_23, Schm2_30
484	486	G	D	Schm1_17, Schm1_92, Schm1_144, Schm1_194, Schm1_234, Schm1_22, Schm1_97, RDN02
493	495	P	L	RDN120
512	514	V	L	Schm1_253, Schm1_123
519	521	Р	S	Schm1_253, Schm1_123
530	532	Α	S	Schm1_141, Schm1_156, Schm1_174
535	537	1	V	RDN120
547	549	Α	V	Schm1_35, Schm1_41, Schm2_9
553	555	G	Т	RDN116
560	562	E	V	RDN02, RDN116
630	632	V	1	RDN75
668	670	T	M	RDN116
689	691	G	Đ	Schm1_39, Schm1_248, Schm1_59, Schm2_14
706	708		V	RDN02
723	725			Schm1_39, Schm1_55, Schm1_56, Schm1_92, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_174, Schm1_234, Schm1_248, Schm1_248, Schm1_29, Schm1_41, Schm1_59, Schm1_41, Schm1_59, Schm1_103, Schm1_103, Schm1_103, Schm1_136, Schm1_136, Schm2_9, Schm2_30, Schm2_30, Schm2_50, RDN60,

RDN02, RDN78, RDN120,	
ן אחם ו	
I I KUNTZU,	Ì
RDN116	
734 736 T A RDN02	
743 745 R H RDN116	
749 751 H R Schm1_17,	
Schm1_39,	
Schm1_55,	
Schm1_56,	ļ
Schm1_76,	
Schm1_92,	
Schm1_142,	
Schm1_144,	
Schm1_194,	
Schm1_253,	
Schm1_35,	
Schm1_68,	
Schm1_108,	
Schm1_141,	
Schm1_156,	
Schm1_174,	
Schm1_176,	
Schm1_177,	
Schm1_234,	
Schm1_248,	
Schm1_240, Schm1_3,	
Schm1_22,	
Schm1_25,	
Schm1_29,	
Schm1_41,	
Schm1_43,	
Schm1_59,	
Schm1_85,	
Schm1_97,	
Schm1_99,	
Schm1_103,	
Schm1_123,	
Schm1_136,	
Schm2_9,	
Schm2_3, Schm2_14,	
Schm2_14, Schm2_23,	
Schm2_30,	
Schm2_32,	
Schm2_46,	
RDN60,	
RDN02,	
RDN136,	
RDN78,	
RDN120,	
RDN75,	
RDN116	
770 772 R K RDN60,	
RDN120	
804 806 D A Schm1_55,	
Schm1_68,	
Schm1_248,	
Schm1_3,	
Schm1_3, Schm1_29,	
Schm2_23,	
Schm2_30,	
RDN02,	
RDN120,	
RDN75	
874 876 T M Schm1_35,	

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				Schm1_55, Schm1_56, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_108, Schm1_1141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_234, Schm1_248, Schm1_22, Schm1_25, Schm1_25, Schm1_25, Schm1_99, Schm1_99, Schm1_99, Schm1_103, Schm1_99, Schm1_103, Schm1_123, Schm2_124, Schm2_124, Schm2_23, Schm2_30, Schm2_32, Schm2_32, Schm2_32, Schm2_32,	
				RDN02, RDN136, RDN78, RDN120,	
				RDN75, RDN116	
1241	1243	1	V	Schm1_253,	
1302	1304	D	G	Schm1_123 Schm1_253,	
				Schm1_123	
1313	1315	D	G	Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_92, Schm1_94, Schm1_142, Schm1_144, Schm1_1253, Schm1_12X, Schm1_12X, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156,	

				Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_5, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_99, Schm1_103, Schm1_103, Schm1_136, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_50,
				RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
1322	1324	V	1	RDN120
1349	1351	V	M	RDN02
1355	1357	P	S	Schm1_234, Schm1_136, RDN75
1364	1366	R	E	Schm1_156
1365	1367	D	1	Schm1_156
1393	1395	A	V	Schm1_35, Schm1_41, Schm2_9, RDN78
1425	1427	Α	V	RDN02
1479	1481	N	K	RDN60
1483	1485	V	1	Schm1_141, Schm1_156, Schm1_174
1487	1489		M	Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_35, Schm1_36, Schm1_108, Schm1_141, Schm1_141,

				Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_123, Schm1_136, Schm2_14, Schm2_14, Schm2_30, Schm2_32, Schm2_30, Schm2_32, Schm2_30, Schm2_30, Schm2_32, Schm2_30, RDN136, RDN136, RDN136, RDN136, RDN78, RDN120, RDN75, RDN116	
1505 1516	1507 1518	E D	K G	Schm2_50 Schm1_17,	
				Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_92, Schm1_142, Schm1_194, Schm1_194, Schm1_253, Schm1_68, Schm1_108, Schm1_174, Schm1_176, Schm1_176, Schm1_177, Schm1_177, Schm1_177, Schm1_234, Schm1_248, Schm1_248, Schm1_25, Schm1_29, Schm1_29, Schm1_41, Schm1_41, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_97, Schm1_99, Schm1_103, Schm1_103, Schm1_103, Schm1_103, Schm1_103, Schm1_103, Schm1_103, Schm1_103, Schm1_103, Schm1_103,	

1522	1524	E	G	Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_99, Schm2_32,
1538	1540	G	D	Schm2_46 Schm1_17,
1556	1040			Schm1_22, Schm1_97
1545	1547	S	Т	Schm2_50
1555	1557	N	D	Schm1_35, Schm1_41, Schm2_9, RDN78
1560	1562	Τ	A	Schm1_17, Schm1_144, Schm1_194, Schm1_35, Schm1_234, Schm1_22, Schm1_41, Schm1_97, Schm1_99, Schm1_103, Schm1_136, Schm2_9, Schm2_32, Schm2_46, RDN78
1576	1578	G	R	Schm2_50
1580	1582	D	G	Schm1_144, Schm1_194, Schm1_234, Schm1_136
1587	1589	V	Α	Schm1_142, Schm1_176, Schm1_25
1591	1593	N	S	RDN75
1598	1600	Α	V	Schm1_17, Schm1_22, Schm1_97
1605	1607	S	T	Schm1_17, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_35, Schm1_68, Schm1_108,

				Schm1_141, Schm1_176, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_97, Schm1_103, Schm1_103, Schm1_123, Schm1_123, Schm1_123, Schm1_136, Schm2_9, Schm2_14, Schm2_23, Schm2_30, Schm2_32, Schm2_30, Schm2_30, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
1608	1610	S	Р	Schm1_144, Schm1_194, Schm1_234, Schm1_136
1609	1611	Α	Deletion	Schm1_142, Schm1_176, Schm1_25, RDN120
1610	1612	T	Deletion	Schm1_142, Schm1_176, Schm1_25, RDN120
1617	1619	T	A	Schm1_17, Schm1_39, Schm1_56, Schm1_92, Schm1_35, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_248, Schm1_22, Schm1_41, Schm1_59, Schm1_59, Schm1_97, Schm1_97, Schm1_99, Schm2_9, Schm2_14, Schm2_14, Schm2_13,

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				Schm2_30, Schm2_46, Schm2_50, RDN60, RDN78, RDN116
1622	1624	G	S	Schm1_142, Schm1_176, Schm1_25, RDN120
1642	1644	K	T	Schm1_144

Sequence analyses of Spy0488

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Sequences were obtained from all 51 strains. The level of amino acid sequence identity ranged from 85.4% to 100% as compared to the sequence of Spy0488 from *S. pyogenes* SF370. Table 8 lists all 49 amino acid positions which showed a distinct amino acid as compared to Spy0488 from *S. pyogenes* SF370. The genes from several strains (e.g. Schmitz 1/55) possessed furthermore a different N terminus, with an addition of 25 amino acids and a frame-shift for the first 16 amino acids relative to Spy0488 from *S. pyogenes* SF370.

Table 8: Gene conservation of Spy0488. ¹, observed amino acid at respective position in any of the sequenced genes of the respective *S. pyogenes* strains. ², second possible amino acid observed at the respective position. Insertion refers to an additional amino acid relative to Spy0488 of *S. pyogenes* SF370.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	AA change ²	Strains with respective change ¹	Strains with respective change ²
Insertion	1		M		Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_22, Schm1_25, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_59, Schm1_59, Schm1_59,	

1	ı			Schm1_97,
				Schm1_99,
				Schm1_103, Schm1_123,
				Schm1_136,
				Schm2_9,
				Schm2_23, Schm2_30,
1				Schm2_32,
				Schm2_46,
		ļ		Schm2_50,
				RDN60,
İ	i		Į.	RDN02,
				RDN136,
			į l	RDN78,
				RDN120,
				RDN75,
la a antia a	2		NA .	RDN116
Insertion	2	-	M	Schm1_39, Schm1_55,
				Schm1_56,
				Schm1_74,
)	Schm1_76,
				Schm1_92,
			1	Schm1_142, Schm1_144,
				Schm1_194, Schm1_253,
				Schm1_35,
				Schm1_68,
	·			Schm1_108, Schm1_141, Schm1_156, Schm1_174,
				Schm1_176, Schm1_177,
	ļ	ļ		Schm1_234, Schm1_248,
				Schm1_3,
1		l	ł	Schm1_22,
				Schm1_25,
				Schm1_29,
]]		Schm1_41,
				Schm1_43,
)		Schm1_59, Schm1_85,
				Schm1_97,
		}		Schm1_99,
				Schm1_103, Schm1_123,
				Schm1_136,
				Schm2_9,
				Schm2_23,
				Schm2_30,
1		L		Schm2_32,
				Schm2_46, Schm2_50,
				RDN60,
				RDN02,
				RDN136,
				RDN78,
				RDN120,
				RDN75,
			B.78	RDN116
Insertion	3	-	M	Schm1_39, Schm1_55, Schm1_56, Schm1_74,
				Schm1_56, Schm1_74, Schm1_76, Schm1_92,
				Schm1_142, Schm1_144,
				Schm1_194, Schm1_253,
	}			Schm1_35, Schm1_68,
				Schm1_108, Schm1_141,
				Schm1_156, Schm1_174,
		<u> </u>		Schm1_176, Schm1_177,

1	1	1	1	Schm1_234, Schm1_248,
				Schm1_3, Schm1_22,
ļ (1	Schm1_25, Schm1_29,
				Schm1_23, Schm1_23, Schm1_41, Schm1_43,
			į	
1		Ì		Schm1_59, Schm1_85,
			ĺ	Schm1_97, Schm1_99,
Į į	1	ļ	ļ	Schm1_103, Schm1_123,
				Schm1_136, Schm2_9,
				Schm2_23, Schm2_30,
1	İ	- 1	ţ	Schm2_32, Schm2_46,
]			İ	Schm2 50, RDN60,
				RDN02, RDN136, RDN78,
1		1	1	
				RDN120, RDN75, RDN116
Insertion	4	-	L	Schm1_39, Schm1_55,
				Schm1_56, Schm1_74,
1				Schm1 76, Schm1 92,
1	1	-		Schm1 142, Schm1 144,
				Schm1 194, Schm1_253,
	1			_
1	ì	1		Schm1_35, Schm1_68,
		- 1		Schm1_108, Schm1_141,
[Schm1_156, Schm1_174,
]]	1]	Schm1_176, Schm1_177,
			l	Schm1_234, Schm1_248,
1		ļ	ļ	Schm1_3, Schm1_22,
	İ			Schm1_25, Schm1_29,
]		i		Schm1_41, Schm1_43,
1	1			
				Schm1_59, Schm1_85,
				Schm1_97, Schm1_99,
				Schm1_103, Schm1_123,
				Schm1_136, Schm2_9,
				Schm2_23, Schm2_30,
				Schm2_32, Schm2_46,
1				
1				
				Schm2_50, RDN60,
				Schm2_50, RDN60, RDN02, RDN136, RDN78,
				Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_1253,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_1253, Schm1_195, Schm1_68,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_1253, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_1253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_1253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_1253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_3, Schm1_29, Schm1_41, Schm1_43,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_234, Schm1_248, Schm1_25, Schm1_29, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123,
Insertion	5	-	R	Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_99, Schm1_103, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9,
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Insertion	16			Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_32, Schm2_30, Schm2_32, Schm2_30, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
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Insertion	18	-	С	Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141,

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Insertion	19		A	Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_23, Schm2_30, Schm2_32, Schm2_30, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
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Insertion	21.	-	Т	Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68,

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Insertion	23			Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_23, Schm2_30, Schm2_32, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
Insertion	24	-	V	Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253,

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Insertion	25		S		Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_3, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_59, Schm1_85, Schm1_103, Schm1_123, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_23, Schm2_30, Schm2_32, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116	
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				Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9,
				Schm2_23, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
16	41	Α	R	Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_23, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43,
				Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_23, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
30	55	S	F	Schm1_99, Schm1_136, Schm2_46
35	60	S	Υ	RDN75
53	75	A Z	D	Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm2_9, Schm2_23, Schm2_30, Schm2_32, Schm2_30, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116 Schm1_253, Schm1_99,
53	78	N		Schm1_253, Schm1_99, Schm1_123, Schm1_136,

			,	
				Schm2_46, RDN120
56	81	S (Y	Schm1_39, Schm1_55,
				Schm1_56, Schm1_76,
				Schm1_92, Schm1_144,
}	Ì	ì		Schm1_194, Schm1_68,
				Schm1_108, Schm1_141,
1		į		Schm1_156, Schm1_174,
				Schm1_177, Schm1_234,
				Schm1_248, Schm1_3,
)	Ì	ì		Schm1_22, Schm1_29,
				Schm1_43, Schm1_59,
	l l			Schm1_85, Schm1_97,
				Schm1_99, Schm1_103,
				Schm1_136, Schm2_23,
				Schm2_30, Schm2_32,
				Schm2_46, Schm2_50,
İ				RDN60, RDN02, RDN136,
				RDN120, RDN75, RDN116
60	85	D	G	Schm1 248, Schm1 59
69	94	D	G	Schm1_39, Schm1_55,
				Schm1_56, Schm1_76,
}				Schm1_92, Schm1_253,
				Schm1_68, Schm1_108,
				Schm1_00, Cchm1_100, Schm1_141, Schm1_156,
				Schm1_174, Schm1_177,
				Schm1_174, Schm1_17, Schm1_3,
]		<u>'</u>		Schm1_22, Schm1_29,
				Schm1_43, Schm1_59,
,				Schm1_43, Schm1_97,
			•	Schm1_99, Schm1_123,
				Schm1_136, Schm2_23,
				Schm2_30, Schm2_32,
				Schm2_46, Schm2_50,
				RDN60, RDN02, RDN136,
				RDN120
75	100	Q	Н	Schm2_32
76	101	1	T	Schm1_39, Schm1_55,
}	1	1	}	Schm1_56, Schm1_76,
				Schm1_144, Schm1_194,
ļ	ţ	Į		Schm1_253, Schm1_68,
				Schm1_108, Schm1_141,
		!	ł	Schm1_156, Schm1_174,
]]]	Schm1_177, Schm1_234,
1				Schm1_248, Schm1_3,
{		1	\	Schm1_22, Schm1_29,
1				Schm1_43, Schm1_59,
l	l	[(Schm1_85, Schm1_97,
				Schm1_99, Schm1_103,
1				Schm1_123, Schm1_136,
}		1	}	Schm2_23, Schm2_30,
		1		Schm2_32, Schm2_46,
1	[Į.	ļ	Schm2_50, RDN60,
				RDN02, RDN136, RDN120
87	112	F	L	Schm1_253, Schm1_123
93	118	G	E	Schm1 99, Schm2 46
112	137	V	A	Schm1 253, Schm1 123
117	142	i	T	Schm1_39, Schm1_55,
'''	172	'	1	Schm1_56, Schm1_74,
l			Į	Schm1_76, Schm1_92,
]		Schm1_253, Schm1_35,
				Schm1_68, Schm1_108,
}	}	1	1	Schm1_141, Schm1_156,
				Schm1_174, Schm1_177,
l		ļ		Schm1_248, Schm1_3,
	_			
				Schm1_22, Schm1_29,

1 1	1	i		Schm1_41, Schm1_43,
1				Schm1_59, Schm1_85,
1				
				Schm1_97, Schm1_99,
				Schm1_123, Schm1_136,
		į		Schm2_9, Schm2_23,
		1		Schm2 30, Schm2_32,
				Schm2 46, Schm2 50,
		Ì		RDN60, RDN02, RDN136,
				RDN78, RDN120, RDN75,
				RDN116
127	152	H _	Υ	Schm1_39
157	182	D	G	RDN75
163	188	V	1	RDN75
		K	_	Schm1_55, Schm1_68,
174	199	^	ı	
				Schm1_3, Schm1_29,
				Schm2_23, Schm2_30
183	208	G	R	RDN75
184	209	G	S	Schm1_56, Schm1_108,
	200		•	Schm1_85, Schm2_50,
				RDN02
1.5.5	0.10			
188	213	F	L	Schm1_92, Schm1_144,
				Schm1_194, Schm1_253,
				Schm1_35, Schm1_234,
1	\	\		Schm1_41, Schm1_99,
				Schm1_103, Schm1_123,
				Schm1_136, Schm2_9,
	Ì	Ì		
				Schm2_46, RDN78
198	223	P	S	Schm1_92
199	224	K	R	Schm1_56, Schm1_108,
				Schm1_85, Schm2_50,
		ļ		RDN02
004	200		-	
201	226	R	G	Schm1_56, Schm1_74,
		1		Schm1_76, Schm1_92,
				Schm1_144, Schm1_194,
				Schm1_253, Schm1_35,
				Schm1_108, Schm1_177,
				Schm1_234, Schm1_41,
				Schm1_43, Schm1_85,
				Schm1_99, Schm1_103,
				Schm1_123, Schm1_136,
] [İ			Schm2_9, Schm2_32,
				Schm2_46, Schm2_50,
1				RDN02, RDN136, RDN78,
				RDN120
202	227	Q	L	Schm1_144, Schm1_194,
				Schm1_35, Schm1_234,
[]				Schm1_41, Schm1_99,
j ì				Schm1_103, Schm1_136,
				Schm2_9, Schm2_46,
1				
				RDN78
206	231	T		Schm1_56, Schm1_108,
				Schm1_85, Schm2_50,
			•	RDN02
209	234	D	Α	Schm1_92, Schm1_144,
209	254			
				Schm1_194, Schm1_35,
				Schm1_234, Schm1_41,
]]	'			Schm1_99, Schm1_103,
1				Schm1_136, Schm2_9,
			l	Schm2_46, RDN78
217	242	Р	S	Schm1_56, Schm1_108,
217	242		٦	
\ \ \			 	Schm1_85, Schm2_50,
				RDN02
221	246	W	С	Schm1_76, Schm1_177,
				Schm1_43, RDN136
222	247	K	Е	Schm1 56, Schm1 108,
	- T 3	<u>``</u>		

				Schm1_85, Schm2_50, RDN02
232	257	А	T	Schm1_39, Schm1_22, Schm1_97
235	260	S	F	Schm1_253, Schm1_123
238	263	T	1	Schm1_248, Schm1_59
258	283	Α	V	Schm1_92
291	316	E	Q	Schm1_55, Schm1_68, Schm1_3, Schm1_29, Schm2_23, Schm2_30

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Sequences were obtained from all 51 strains. The level of amino acid sequence identity ranged from 98.2% to 100% as compared to the sequence of Spy0872 from *S. pyogenes* SF370. Table 9 lists all 34 acid positions which showed a distinct amino acid as compared to Spy0872 from *S. pyogenes* SF370. The gene from strain Schmitz 1/22 showed in addition an insertion of 2 amino acids after position 587.

Table 9: Gene conservation of Spy0872. 1, observed amino acid at respective position in any of the sequenced genes of the respective *S. pyogenes* strains. Insertion refers to an additional amino acid relative to Spy0872 of *S. pyogenes* SF370.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	Strains with respective change
67	67	G	С	Schm1_136
74	74	E	D	Schm1_76, Schm1_177, Schm1_43, RDN136
178	178	K	N	Schm1_7, Schm1_39, Schm1_55, Schm1_56, Schm1_74, Schm1_76, Schm1_92, Schm1_142, Schm1_144, Schm1_194, Schm1_253, Schm1_35, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_41, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm1_123, Schm1_136, Schm1_99, Schm1_14, Schm2_32, Schm1_136, Schm2_9, Schm2_14, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN02, RDN136, RDN78, RDN120, RDN75, RDN116
181	181	Р	S	RDN60
222	222	Н	Y	RDN120
228	228	V	Α	Schm1_56, Schm1_108, Schm1_85, Schm2_50
253	253	V		Schm1_7, Schm1_39, Schm1_55, Schm1_56, Schm1_76, Schm1_142, Schm1_144, Schm1_194, Schm1_68, Schm1_108, Schm1_141, Schm1_156, Schm1_174, Schm1_176, Schm1_177, Schm1_234, Schm1_248, Schm1_3, Schm1_22, Schm1_25, Schm1_29, Schm1_43, Schm1_59, Schm1_85, Schm1_97, Schm1_99, Schm1_103, Schm2_14, Schm2_32, Schm2_46, Schm2_50, RDN60, RDN136, RDN120
328	328	Ī	M	Schm1_55, Schm1_56, Schm1_92, Schm1_68, Schm1_108, Schm1_3, Schm1_29, Schm1_85, Schm1_136, Schm2_23, Schm2_30, Schm2_50, RDN75

			_	
329	329	K	Т	Schm1_55, Schm1_56, Schm1_92, Schm1_68,
				Schm1_108, Schm1_3, Schm1_29, Schm1_85,
				Schm1_136, Schm2_23, Schm2_30, Schm2_50, RDN75
336	336	V	1	Schm1_56, Schm1_108, Schm1_85, Schm2_50
337	337	Α	T	Schm1_136, RDN75
340	340	P	L	RDN120
393	393	A	V	Schm1_7, Schm1_39, Schm1_55, Schm1_56,
				Schm1_74, Schm1_76, Schm1_92, Schm1_142,
				Schm1_144, Schm1_194, Schm1_253, Schm1_35,
		1		Schm1_68, Schm1_108, Schm1_141, Schm1_156,
				Schm1_174, Schm1_176, Schm1_177, Schm1_234,
		\		Schm1_248, Schm1_3, Schm1_22, Schm1_25,
				Schm1_29, Schm1_41, Schm1_43, Schm1_59,
				Schm1_85, Schm1_97, Schm1_99, Schm1_103,
				Schm1_123, Schm1_136, Schm2_9, Schm2_14,
				Schm2_23, Schm2_30, Schm2_32, Schm2_46,
1		1		Schm2_50, RDN60, RDN02, RDN136, RDN78,
l i				RDN120, RDN75, RDN116
412	412	M	1	RDN120
427	427	D	Υ	Schm2_46
433	433	G	E	Schm1_7, Schm1_22, Schm1_97
444	444		Т	RDN75
478	478	Y	F	Schm1_253, Schm1_123
490	490	T	1	Schm1_55, Schm1_68, Schm1_3, Schm1_29,
				Schm2_23, Schm2_30
492	492	F	С	RDN02
532	532	Α	T	Schm1_144, Schm1_194, Schm1_234, Schm1_103
535	535		V	Schm1_142, Schm1_176, Schm1_25, Schm2_46,
				RDN116
553	553	E	Q	Schm1_142, Schm1_176, Schm1_25, Schm1_99,
				Schm2_32, Schm2_46, RDN116
576	576	S	R	Schm1_142, Schm1_176, Schm1_25, Schm1_99,
				Schm2_46, RDN116
580	580	V	ı	Schm1_142, Schm1_176, Schm1_25, Schm1_99,
				Schm2_46, RDN116
Insertion	588	-		Schm1_7, Schm1_22, Schm1_97
Insertion	589	-	l	Schm1_7, Schm1_22, Schm1_97
588	590	1	T	RDN78
598	600	G	D	Schm1_92
600	602) T	1	Schm1_7, Schm1_39, Schm1_55, Schm1_56,
				Schm1_74, Schm1_76, Schm1_92, Schm1_142,
1		1		Schm1_144, Schm1_194, Schm1_253, Schm1_35,
				Schm1_68, Schm1_108, Schm1_141, Schm1_156,
1		1		Schm1_174, Schm1_176, Schm1_177, Schm1_234,
				Schm1_248, Schm1_3, Schm1_22, Schm1_25,
				Schm1_29, Schm1_41, Schm1_43, Schm1_59,
		1		Schm1_85, Schm1_97, Schm1_99, Schm1_103,
				Schm1_123, Schm1_136, Schm2_9, Schm2_14,
1				Schm2_23, Schm2_30, Schm2_32, Schm2_46,
				Schm2_50, RDN60, RDN02, RDN136, RDN78,
				RDN120, RDN75, RDN116
605	607	\ \ \		Schm1_7, Schm1_39, Schm1_56, Schm1_76,
				Schm1_144, Schm1_194, Schm1_253, Schm1_108,
1				Schm1_141, Schm1_156, Schm1_174, Schm1_177,
				Schm1_234, Schm1_248, Schm1_22, Schm1_43,
-				Schm1_59, Schm1_85, Schm1_97, Schm1_103,
				Schm1_123, Schm2_14, Schm2_50, RDN60, RDN02,
222	000	 		RDN136, RDN78, RDN120
620	622	L	F	Schm1_7, Schm1_142, Schm1_176, Schm1_22,
				Schm1_25, Schm1_97, Schm1_99, Schm2_32,
605	607	+ -	1	Schm2_46, RDN116
625	627	S	N	Schm1_7, Schm1_22, Schm1_97 Schm1_7, Schm1_142, Schm1_176, Schm1_22,
634	636	3	in in	Schm1_7, Schm1_142, Schm1_176, Schm1_22, Schm1_25, Schm1_97, Schm1_99, Schm2_46, RDN116_
			L	John 11_20, John 11_37, John 11_30, John 12_40, Kolario

659	661	G	С	Schm1_253, Schm1_123
667	669	К	Ε	Schm1_144, Schm1_194, Schm1_234, Schm1_103, RDN120

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Sequences were obtained from all 51 strains. The level of amino acid sequence identity ranged from 98.9% to 100% as compared to the sequence of Spy0895 from *S. pyogenes* SF370. Table 10 lists all 13 amino acid positions which showed a distinct amino acid as compared to Spy0895 from *S. pyogenes* SF370.

Table 10: Gene conservation of Spy0895. 1, observed amino acid at respective position in any of the sequenced genes of the respective S. pyogenes strains.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	Strains with respective change
19	19	Α	V	Schm1_17, Schm1_22, Schm1_97
33	33	Α	V	Schm1_17, Schm1_141, Schm1_156, Schm1_174, Schm1_22, Schm1_97, RDN02
50	50	F	V	Schm1_253, Schm1_123
52	52	Α	V	Schm1_17, Schm1_55, Schm1_68, Schm1_141, Schm1_156, Schm1_174, Schm1_3, Schm1_22, Schm1_29, Schm1_97, Schm2_30
60	60	Т	1	Schm1_56, Schm1_108, Schm1_85, Schm2_50
71	71	L	ı	Schm1_92, Schm1_144, Schm1_194, Schm1_234, Schm1_103
138	138	Н	Q	Schm1_92, Schm1_144, Schm1_194, Schm1_234, Schm1_103
188	188	R	P	Schm1_174
238	238	R	С	Schm1_55, Schm1_76, Schm1_68, Schm1_177, Schm1_3, Schm1_29, Schm1_43, Schm2_30, RDN136
242	242	Y	С	Schm1_136
252	252	K	Т	Schm1_56, Schm1_108, Schm1_85, Schm2_50
255	255	S	G	Schm1_56, Schm1_108, Schm1_85, Schm2_50
256	256	L	F	RDN60

Sequence analyses of Spy1536

Sequences were obtained from all 51 strains. The level of amino acid sequence identity ranged from 99.1% to 100% as compared to the sequence of Spy1536 from *S. pyogenes* SF370. Table 11 lists all 8 amino acid positions which showed a distinct amino acid as compared to Spy1536 from *S. pyogenes* SF370. The gene from strain Schmitz 2/14 showed in addition an insertion of 3 amino acids after position 207.

Table 11: Gene conservation of Spy1536. \(^1\), observed amino acid at respective position in any of the sequenced genes of the respective S. pyogenes strains. Insertion refers to an additional amino acid relative to Spy1536 of S. pyogenes SF370.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	Strains with respective change
5	5	K	N	Schm1_12, Schm2_9, Schm1_136
92	92	G	R	Schm1_142
97	97	Α	T	Schm1_5, Schm1_74
125	125	Р	S	Schm1_123
126	126	V	Α	Schm1_142
183	183	V	1	Schm1_94, RDN78, Schm1_97, Schm1_59, Schm1_76, RDN136, Schm1_177, Schm2_32, Schm1_141, Schm1_144, RDN120, Schm1_25, Schm1_176, RDN75_85, Schm2_46, Schm2_23, Schm1_55
Insertion	208	-	K	Schm2_14
Insertion	209	_	N	Schm2_14
Insertion	210	_	G	Schm2_14
333	336	V	1	Schm1_12, Schm1_35, Schm2_9, Schm1_174, Schm1_136, Schm1_234, Schm1_68
337	340	Q	E	Schm1_43, Schm1_108

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Sequences were obtained from 50 strains. The sequence from strain RDN120 was not determined. The level of amino acid sequence identity ranged from 98.2 to 100% as compared to the sequence of Spy1666 from *S. pyogenes* SF370. Table 12 lists all 18 amino acid positions which showed a distinct amino acid as compared to Spy1666 from *S. pyogenes* SF370.

Table 12: Gene conservation of Spy1666. , observed amino acid at respective position in any of the sequenced genes of the respective S. pyogenes strains.

Position in SF370 gene	Alignment position	Amino acid in SF370 gene	AA change ¹	Strains with respective change
3	3	S	Р	Schm1_17, Schm1_22, Schm1_97, Schm1_136, Schm1_17, Schm1_22, Schm1_97, Schm1_136
11	11	L	V	Schm1_17, Schm1_22, Schm1_97, Schm1_136, Schm1_17, Schm1_22, Schm1_97, Schm1_136
45	45	D	N	Schm1_17, Schm1_22, Schm1_97, Schm1_136, Schm1_17, Schm1_22, Schm1_97, Schm1_136
67	67	G	S	Schm1_17, Schm1_22, Schm1_97, Schm1_136, Schm1_17, Schm1_22, Schm1_97, Schm1_136
69	69	E	Q	Schm1_17, Schm1_22, Schm1_97, Schm1_136, Schm1_17, Schm1_22, Schm1_97, Schm1_136
90	90	К	Q	Schm1_142, Schm1_176, Schm1_25, Schm2_46, Schm1_142, Schm1_176, Schm1_25, Schm2_46
106	106	R		RDN136, RDN78, RDN136, RDN78

120	120		F_	Schm1_136, Schm1_136
149	149	L	S	RDN78, RDN78
167	167	T	N	RDN75, RDN75
204	204	Т	Α	Schm1_253, Schm1_103, Schm1_123, Schm1_253, Schm1_103, Schm1_123
217	217	Р	S	Schm1_39, Schm1_248, Schm1_59, Schm1_39, Schm1_248, Schm1_59
251	251	Q	Н	Schm1_97, Schm1_97
252	252	D	E	Schm1_76, Schm1_141, Schm1_156, Schm1_174, Schm1_177, Schm1_43, Schm2_32, RDN136, Schm1_76, Schm1_141, Schm1_156, Schm1_174, Schm1_177, Schm1_43, Schm2_32, RDN136
259	259	L	F	Schm1_92, RDN75, Schm1_92, RDN75
292	292	L	F	RDN116, RDN116
302	302	К	Т	Schm1_17, Schm1_142, Schm1_176, Schm1_22, Schm1_25, Schm1_97, Schm2_46, Schm1_17, Schm1_142, Schm1_176, Schm1_22, Schm1_25, Schm1_97, Schm2_46
319	319	Ť	Α	Schm1_76, Schm1_141, Schm1_156, Schm1_174, Schm1_177, Schm1_43, Schm2_32, RDN136, Schm1_76, Schm1_141, Schm1_156, Schm1_174, Schm1_177, Schm1_43, Schm2_32, RDN136

No sequence variation was observed on the amino acid sequence level in any of the analyzed 51 gene sequences obtained from the listed *S. pyogenes* strains.

SEQUENCE DATA FOR AMINO ACID SEQUENCES

1. Spy0269

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1.1 Full length Spy0269

> Spy0269 / SF370 (serotype 1); SEQ ID NO: 57

MDLEQTKPNQVKQKIALTSTIALLSASVGVSHQVKADDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKT
LSQQKAELTELATALTKTTAEINHLKEQQDNEQKALTSAQEIYTNTLASSEETLLAQGAEHQRELTATETELH
NAQADQHSKETALSEQKASISAETTRAQDLVEQVKTSEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELE
KAKADLENQKAKVKKQLTEELAAQKAALAEKEAELSRLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASG
YIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPADRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLP
PVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPGVSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGA
FNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAINFLRVDKHNPNAPVYLGFSTSNVGSLNEHFVMFPESN
IANHQRFNKTPIKAVGSTKDYAQRVGTVSDTIAAIKGKVSSLENRLSAIHQEADIMAAQAKVSQLQGKLASTL
KQSDSLNLQVRQLNDTKGSLRTELLAAKAKQAQLEATRDQSLAKLASLKAALHQTEALAEQAAARVTALVAKK
AHLQYLRDFKLNPNRLQVIRERIDNTKQDLAKTTSSLLNAQEALAALQAKQSSLEATIATTEHQLTLLKTLAN
EKEYRHLDEDIATVPDLQVAPPLTGVKPLSYSKIDTTPLVQEMVKETKQLLEASARLAAENTSLVAEALVGQT
SEMVASNAIVSKITSSITQPSSKTSYGSGSSTTSNLISDVDESTQRALKAGVVMLAAVGLTGFRFRKESK

20 1.2 Antigenic fragment Spy0269-1

> Spy0269-1 / SF370 (serotype 1); SEQ ID NO: 1
DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL
TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
NFLRVDKHNPNAPV

1.3 Homologous sequences of other S. pyogenes isolates and/or serotypes

> Spy0269-1 / Schmitz 2/14 (serotype 1); SEQ ID NO: 58
DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINNLKEQQDNEQKAL

TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
NFLRVDKRNPNAPV

> Spy0269-1 / Schmitz 1/156 (serotype 4); SEQ ID NO: 59
DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL
TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
NFLRVDKHNPKAPV

> Spy0269-1 / Schmitz 1/59 (serotype 12); SEQ ID NO: 60
DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTKLATALTKTTAEINHLKEQQDNEQKAL
TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
NFLRVDKRNPNAPV

> Spy0269-1 / Schmitz 1/177 (serotype 22); SEQ ID NO: 61
DDRASGETKASNTHDDSLPKPETIQEAKATIEAVEKALSQQKAELTELATALTKTTAKINHLKEQQDNEQKAL
TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
NFLRVDKRNPNAPV

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- > Spy0269-1 / Schmitz 1/136 (serotype 25); SEQ ID NO: 63

 DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL
 TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
 SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
 RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
 DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
 VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
 NFLRVDKRNPNAPV
- > Spy0269-1 / Schmitz 1/85 (serotype 28); SEQ ID NO: 64
 DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL

 TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
 SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
 RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
 DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
 VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
 NFLRVDKHNPNAPV
 - > Spy0269-1 / Schmitz 2/50 (serotype 28); SEQ ID NO: 65
 DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL
 TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQVDQHSKETALSEQKASISAETTRAQDLVEQVKT
 SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
 RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
 DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
 VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
 NFLRVDKRNPNAPV
- > Spy0269-1 / Schmitz 1/123 (serotype 49); SEQ ID NO: 66
 DDRASGETKASNTHDDSLPKPETIQEAKATIDAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL
 TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT
 SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS
 RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA
 DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG
 VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI
 NFLRVDKRNPNAPV
- > Spy0269-1 / Schmitz 1/176 (serotype 83); SEQ ID NO: 67 DDRASGETKASNTHDDSLPKPETIQEAKATIEAVEKTLSQQKAELTELATALTKTTAEINHLKEQQDNEQKAL TSAQEIYTNTLASSEETLLAQGAEHQRELTATETELHNAQADQHSKETALSEQKASISAETTRAQDLVEQVKT SEQNIAKLNAMISNPDAITKAAQTANDNTKALSSELEKAKADLENQKAKVKKQLTEELAAQKAALAEKEAELS RLKSSAPSTQDSIVGNNTMKAPQGYPLEELKKLEASGYIGSASYNNYYKEHADQIIAKASPGNQLNQYQDIPA DRNRFVDPDNLTPEVQNELAQFAAHMINSVRRQLGLPPVTVTAGSQEFARLLSTSYKKTHGNTRPSFVYGQPG

VSGHYGVGPHDKTIIEDSAGASGLIRNDDNMYENIGAFNDVHTVNGIKRGIYDSIKYMLFTDHLHGNTYGHAI NFLRVDKRNPNAPV

2. Spy0292

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2.1 Full length Spy0292

> Spy0292 / SF370 (serotype 1); SEQ ID NO: 68
MIKRLISLVVIALFFAASTVSGEEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKG
KLNWDSPVTISNYPYELTTNYTISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQL
RQWGISDAKVVNSTGLTNHFLGANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFAGQTIY
SYNYMLKGMPCYREGVDGLFVGYSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLIN
FQKVQLIENNKPVKTLYVLDSPEKTVKLVAQNSLFFIKPIHTKTKNTVHITKKSSTMIAPLSKGQVLGRATLQ
DKHLIGQGYLDTPPSINLILQKNISKSFFLKVWWNRFVRYVNTSL

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2.2 Antigenic fragment Spy0292-1

> Spy0292-1 / SF370 (serotype 1); SEQ ID NO: 2
EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
ANTYPNTEPDDENCFC

2.3 Homologous sequences of other S. pyogenes isolates and/or serotypes

- > Spy0292-1 / Schmitz 1/56 (serotype 28); SEQ ID NO: 71
 EEYSVTAKHAIAVDLESGKVLYEKDTKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFC
- > Spy0292-1 / Schmitz 1/74 (serotype 3); SEQ ID NO: 72
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFC
- > Spy0292-1 / Schmitz 1/76 (serotype 22); SEQ ID NO: 73
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFC
- > Spy0292-1 / Schmitz 1/92 (serotype 11); SEQ ID NO: 74
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFC
- > Spy0292-1 / Schmitz 1/94 (serotype 1); SEQ ID NO: 75 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG ANTYPNTEPDDENCFC
- 60 > Spy0292-1 / Schmitz 1/142 (serotype 83); SEQ ID NO: 76

EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG ANTYPNTEPDDENCFC

- 5 > Spy0292-1 / Schmitz 1/144 (serotype 76); SEQ ID NO: 77 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG ANTYPNTEPDDENCFC
- > Spy0292-1 / Schmitz 1/194 (serotype 44); SEQ ID NO: 78 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG ANTYPNTEPDDENCFC

15 2.4 Antigenic fragment Spy0292-3

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> Spy0292-3 / SF370 (serotype 1); SEQ ID NO: 3
EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFAGQTIYSYNYMLKGMPCYREGVDGLFVG
YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE

2.5 Homologous sequences of other S. pyogenes isolates and/or serotypes

- > Spy0292-3 / Schmitz 1/39 (serotype 12); SEQ ID NO: 79 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFVG YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
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 > Spy0292-3 / Schmitz 1/55 (serotype 118); SEQ ID NO: 80
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDTKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/56 (serotype 28); SEQ ID NO: 81
 EEYSVTAKHAIAVDLESGKVLYEKDTKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 40 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/74 (serotype 3); SEQ ID NO: 82
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFAGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/76 (serotype 22); SEQ ID NO: 83

 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/92 (serotype 11); SEQ ID NO: 84
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFIG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE

- > Spy0292-3 / Schmitz 1/94 (serotype 1); SEQ ID NO: 85
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFAGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/142 (serotype 83); SEQ ID NO: 86
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/144 (serotype 76); SEQ ID NO: 87
 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFDGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVINADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE
- > Spy0292-3 / Schmitz 1/194 (serotype 44); SEQ ID NO: 88

 EEYSVTAKHAIAVDLESGKVLYEKDAKEVVPVASVSKLLTTYLVYKEVSKGKLNWDSPVTISNYPYELTTNYT
 ISNVPLDKRKYTVKELLSALVVNNANSPAIALAEKIGGTEPKFVDKMKKQLRQWGISDAKVVNSTGLTNHFLG
 ANTYPNTEPDDENCFCATDLAIIARHLLLEFPEVLKLSSKSSTIFAGQTIYSYNYMLKGMPCYREGVDGLFVG
 YSKKAGASFVATSVENQMRVITVVLNADQSHEDDLAIFKTTNQLLQYLLINFQKVQLIE

25 3. Spy0416A

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3.1 Full length Spy0416A

> Spy0416A / SF370 (serotype 1); SEQ ID NO: 89 ADELSTMSEPTITNHAQQQAQHLTNTELSSAESKSQDTSQITLKTNREKEQSQDLVSEPTTTELADTDAASMA 30 NTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDIMGSAE SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE 35 STDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGAGLLNIDGAVTSGL YVTGKDNYGSISLGNITDTMTFDVTVHNLSNKDKTLRYDTELLTDHVDPQKGRFTLTSHSLKTYQGGEVTVPA NGKVTVRVTMDVSQFTKELTKQMPNGYYLEGFVRFRDSQDDQLNRVNIPFVGFKGQFENLAVAEESIYRLKSQ 40 GKTGFYFDESGPKDDIYVGKHFTGLVTLGSE

3.2 Antigenic fragment Spy0416A-1

45 > Spy0416A-1 / SF370 (serotype 1); SEQ ID NO: 4
 ADELSTMSEPTITNHAQQQAQHLTNTELSSAESKSQDTSQITLKTNREKEQSQDLVSEPTTTELADTDAASMA
 NTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET
 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDIMGSAE
50 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF
 GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS
 MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA

3.3 Homologous sequences of other S. pyogenes isolates and/or serotypes

> Spy0416A-1 / Schmitz 1/7 (serotype 4); SEQ ID NO: 90
ADELTTTSEPTITNHAQQQAQHLTNTELSSAESQSPDTSQITPKTNREKEQPQGLVSEPTTTELADTDAASMA
NTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA

RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFGDFDEDWENFEFDAEPKAIKKNKIYRPQSTQAPKETVI KTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESL FIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYG LVGSPSTGRTPTSVAAINSKWVIQRLMTAKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKEST DAGYKAQDVKDKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGK AMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTSMA SPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA

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- > Spy0416A-1 / Schmitz 1/39 (serotype 12); SEQ ID NO: 91
 ADELTTTSEPTITNHTQQQAQHLTNTELSSAESKPQDTSQITLKTNREKEQPQGLVSEPTTTELADTDAAPMA
 NTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET
 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAE
 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF
 GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS
 MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA
- 20 > Spy0416A-1 / Schmitz 1/55 (serotype 118); SEQ ID NO: 92
 ADELTTTSEPTITNHAQQQAPPLTNTELSSAESQPQDTSQVTPETNREKEQPQGLVSEPTTTELADTDAAPMA
 NTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET
 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAE
 25 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYNAQNVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF
 GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS
 MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA
- > Spy0416A-1 / Schmitz 1/56 (serotype 28); SEQ ID NO: 93
 ADELTTTSEPTITNHAQQQAPPLTNTELSSAESQPQDTSQVTPETNREKEQPQGLVSEPTTTELADTDAAPMA
 NTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET
 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAE
 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYNAQNVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF
 GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS
 MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA
- > Spy0416A-1 / Schmitz 1/94 (serotype 1); SEQ ID NO: 94
 ADELSTMSEPTITNHAQQQAQHLTNTELSSAESKSQDTSQITLKTNREKEQSQDLVSEPTTTELADTDAASMA
 NTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 45 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET
 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDIMGSAE
 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF
 50 GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS
 MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA
- > Spy0416A-1 / Schmitz 1/253 (serotype 49); SEQ ID NO: 95
 ADELTTTSEPTITNHAQQQAQPLTNTELSSAESQSPDISQVTPETNREKEQPQGLVSEPTTTELADTDAAPMA
 NTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDADAEPKAIKKHKIYRPQSTQAPKET
 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAE
 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKGLENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALGLLIFNNKSGQSNRSMRLTANGMGIPSAFISHEF

GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA

- > Spy0416A-1 / Schmitz 1/174 (serotype 6); SEQ ID NO: 96
 5 ADELTTTSEPTITNHAQQQAQHLTNTELSSAESKPQDTSQITPKTNREKEQSQDLVSEPTTTELADTDAASMA NTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKET VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAE SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKNIKDSLGYDKSHQFAYVKE STDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTSNGMGIPSAFISHEF GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA
- > Spy0416A-1 / Schmitz 1/234 (serotype 44); SEQ ID NO: 98
 ADELSTMSEPTITNHAQQQAQHLTNTELSSAESKSQDTSQITPKTNREKEQSQDLVSEPTTTELADTDAASMA
 NTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA
 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDEDWENFEFDADAEPKAIKKHKIYRPQSTQAPKET

 VIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAE
 SLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPD
 YGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKE
 STDAGYKAQDVKDKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEF
 GKAMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTS
 MASPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA
- > Spy0416A-1 / Schmitz 1/22 (serotype 4); SEQ ID NO: 99

 ADELTTTSEPTITNHAQQQAQHLTNTELSSAESQSPDTSQITPKTNREKEQPQGLVSEPTTTELADTDAASMA
 NTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQGKVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLA

 40 RQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFGDFDEDWENFEFDAEPKAIKKNKIYRPQSTQAPKETVI
 KTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESL
 FIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYG
 LVGSPSTGRTPTSVAAINSKWVIQRLMTAKELENRADLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKEST
 DAGYKAQDVKDKIALIERDPNKTYDEMIALAKKHGALGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGK

 45 AMSQLNGNGTGSLEFDSVVSKAPSQKGNEMNHFSNWGLTSDGYLKPDITAPGGDIYSTYNDNHYGSQTGTSMA
 SPQIAGASLLVKQYLEKTQPNLPKEKIADIVKNLLMSNAQIHVNPETKTTTSPRQQGA

3.4 Antigenic fragment Spy0416A-6

> Spy0416A-6 / SF370 (serotype 1); SEQ ID NO: 5 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED WENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE AAATGERFLGIAPEAQVMFMRVFANDIMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN HGKAIYSESVDFKDIKDSL

3.5 Homologous sequences of other S. pyogenes isolates and/or serotypes

> Spy0416A-6 / Schmitz 1/7 (serotype 4); SEQ ID NO: 100

AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFGDFDED

WENFEFDAEPKAIKKNKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAA ATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKA KKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTAKELENRADLNHG KAIYSESVDFKDIKDSL

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- > Spy0416A-6 / Schmitz 1/39 (serotype 12); SEQ ID NO: 101 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED WENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE AAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN HGKAIYSESVDFKDIKDSL
- > Spy0416A-6 / Schmitz 1/55 (serotype 118); SEQ ID NO: 102
 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED
 WENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE
 AAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE
 KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN
 HGKAIYSESVDFKDIKDSL
- > Spy0416A-6 / Schmitz 1/56 (serotype 28); SEQ ID NO: 103 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED WENFEFDAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAA ATGERFLGIAPETQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKA KKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLNHG KAIYSESVDFKDIKDSL
 - > Spy0416A-6 / Schmitz 1/94 (serotype 1); SEQ ID NO: 104
 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED
 WENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE
 AAATGERFLGIAPEAQVMFMRVFANDIMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE
 KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN
 HGKAIYSESVDFKDIKDSL
- > Spy0416A-6 / Schmitz 1/253 (serotype 49); SEQ ID NO: 105

 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED
 WENFEFDADAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE
 AAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE
 KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKGLENRADLN
 HGKAIYSESVDFKDIKDSL

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- > Spy0416A-6 / Schmitz 1/174 (serotype 6); SEQ ID NO: 106
 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED
 WENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE
 AAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE
 KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN
 HGKAIYSESVDFKNIKDSL
- > Spy0416A-6 / Schmitz 1/176 (serotype 83); SEQ ID NO: 107
 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED
 WENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE
 AAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE
 KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN
 HGKAIYSESVDFKNIKDSL
- > Spy0416A-6 / Schmitz 1/234 (serotype 44); SEQ ID NO: 108 AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDFDED WENFEFDADAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKE AAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIE KAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADLN HGKAIYSESVDFKDIKDSL

> Spy0416A-6 / Schmitz 1/22 (serotype 4); SEQ ID NO: 109
AVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFGDFDED
WENFEFDAEPKAIKKNKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSKEAA
ATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAIEKA
KKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTAKELENRADLNHG
KAIYSESVDFKDIKDSL

3.6 Antigenic fragment Spy0416A-7

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20 3.7 Homologous sequences of other S. pyogenes isolates and/or serotypes

> Spy0416A-7 / Schmitz 1/7 (serotype 4); SEQ ID NO: 110
SQITPKTNREKEQPQGLVSEPTTTELADTDAASMANTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFGDF
DEDWENFEFDAEPKAIKKNKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSK
EAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAI
EKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTAKELENRADL
NHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYKAQDVKDKIALIERDPNKTYDEMIALAKKHGALG
VLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS

> Spy0416A-7 / Schmitz 1/39 (serotype 12); SEQ ID NO: 111
SQITLKTNREKEQPQGLVSEPTTTELADTDAAPMANTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF
DEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGN
SKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLME
AIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRA
DLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGA
LGVLIFNNKPGOSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS

- > Spy0416A-7 / Schmitz 1/55 (serotype 118); SEQ ID NO: 112 SQVTPETNREKEQPQGLVSEPTTTELADTDAAPMANTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF DEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGN SKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLME AIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRA DLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYNAQNVKGKIALIERDPNKTYDEMIALAKKHGA LGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS
- > Spy0416A-7 / Schmitz 1/56 (serotype 28); SEQ ID NO: 113

 SQITPKINREKEQPQGLVSEPTTTELADTDAAPMANTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
 KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF
 DEDWENFEFDAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSK
 EAAATGERFLGIAPETQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAI
 EKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRADL
 NHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGALG
 VLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS
 - > Spy0416A-7 / Schmitz 1/94 (serotype 1); SEQ ID NO: 114
 SQITLKTNREKEQSQDLVSEPTTTELADTDAASMANTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
 KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF

105

DEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGN SKEAAATGERFLGIAPEAQVMFMRVFANDIMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLME AIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRA DLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGA LGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS

- > Spy0416A-7 / Schmitz 1/253 (serotype 49); SEQ ID NO: 115
 SQVTPETNREKEQPQGLVSEPTTTELADTDAAPMANTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
 KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF
 DEDWENFEFDADAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGN
 SKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLME
 AIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKGLENRA
 DLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYNAQDVKGKIALIERDPNKTYDEMIALAKKHGA
 LGLLIFNNKSGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS
- > Spy0416A-7 / Schmitz 1/176 (serotype 83); SEQ ID NO: 117 SQITLKTNREKEQPQGLVSEPTTTELADTDAAPMANTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF DEDWENFEFDAEAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGN SKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLME AIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRA DLNHGKAIYSESVDFKNIKDSLGYDKSHQFAYVKESTDAGYKAQDVKGKIALIERDPNKTYDEMIALAKKHGA LGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS
- > Spy0416A-7 / Schmitz 1/234 (serotype 44); SEQ ID NO: 118

 SQITPKTNREKEQSQDLVSEPTTTELADTDAASMANTGSDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
 KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFEDF
 DEDWENFEFDADAEPKAIKKHKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGN
 SKEAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLME
 AIEKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTVKELENRA
 DLNHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYKAQDVKDKIALIERDPNKTYDEMIALAKKHGA
 LGVLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS
- > Spy0416A-7 / Schmitz 1/22 (serotype 4); SEQ ID NO: 119
 SQITPKTNREKEQPQGLVSEPTTTELADTDAASMANTGPDATQKSASLPPVNTDVHDWVKTKGAWDKGYKGQG
 KVVAVIDTGIDPAHQSMRISDVSTAKVKSKEDMLARQKAAGINYGSWINDKVVFAHNYVENSDNIKENQFGDF
 DEDWENFEFDAEPKAIKKNKIYRPQSTQAPKETVIKTEETDGSHDIDWTQTDDDTKYESHGMHVTGIVAGNSK
 EAAATGERFLGIAPEAQVMFMRVFANDVMGSAESLFIKAIEDAVALGADVINLSLGTANGAQLSGSKPLMEAI
 EKAKKAGVSVVVAAGNERVYGSDHDDPLATNPDYGLVGSPSTGRTPTSVAAINSKWVIQRLMTAKELENRADL
 NHGKAIYSESVDFKDIKDSLGYDKSHQFAYVKESTDAGYKAQDVKDKIALIERDPNKTYDEMIALAKKHGALG
 VLIFNNKPGQSNRSMRLTANGMGIPSAFISHEFGKAMSQLNGNGTGS

3.8 Full length Spy0416B

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> Spy0416B / SF370 (serotype 1); SEQ ID NO: 56
HVDPQKGRFTLTSHSLKTYQGGEVTVPANGKVTVRVTMDVSQFTKELTKQMPNGYYLEGFVRFRDSQDDQLNR
VNIPFVGFKGQFENLAVAEESIYRLKSQGKTGFYFDESGPKDDIYVGKHFTGLVTLGSETNVSTKTISDNGLH
TLGTFKNADGKFILEKNAQGNPVLAISPNGDNNQDFAAFKGVFLRKYQGLKASVYHASDKEHKNPLWVSPESF
KGDKNFNSDIRFAKSTTLLGTAFSGKSLTGAELPDGHYHYVVSYYPDVVGAKRQEMTFDMILDRQKPVLSQAT
FDPETNRFKPEPLKDRGLAGVRKDSVFYLERKDNKPYTVTINDSYKYVSVEDNKTFVERQADGSFILPLDKAK
LGDFYYMVEDFAGNVAIAKLGDHLPQTLGKTPIKLKLTDGNYQTKETLKDNLEMTQSDTGLVTNQAQLAVVHR
NQPQSQLTKMNQDFFISPNEDGNKDFVAFKGLKNNVYNDLTVNVYAKDDHQKQTPIWSSQAGASVSAIESTAW

YGITARGSKVMPGDYQYVVTYRDEHGKEHQKQYTISVNDKKPMITQGRFDTINGVDHFTPDKTKALDSSGIVR EEVFYLAKKNGRKFDVTEGKDGITVSDNKVYIPKNPDGSYTISKRDGVTLSDYYYLVEDRAGNVSFATLRDLK AVGKDKAVVNFGLDLPVPEDKQIVNFTYLVRDADGKPIENLEYYNNSGNSLILPYGKYTVELLTYDTNAAKLE SDKIVSFTLSADNNFQQVTFKITMLATSQITAHFDHLLPEGSRVSLKTAQDQLIPLEQSLYVPKAYGKTVQEG TYEVVVSLPKGYRIEGNTKVNTLPNEVHELSLRLVKVGDASDSTGDHKVMSKNNSQALTASATPTKSTTSATA KALPST

4. Spy0872

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10 4.1 Full length Spy0872

> Spy0872 / SF370 (serotype 1); SEQ ID NO: 120
DQVDVQFLGVNDFHGALDNTGTAYTPSGKIPNAGTAAQLGAYMDDAEIDFKQANQDGTSIRVQAGDMVGASPA
NSALLQDEPTVKVFNKMKFEYGTLGNHEFDEGLDEFNRIMTGQAPDPESTINDITKQYEHEASHQTIVIANVI
DKKTKDIPYGWKPYAIKDIAINDKIVKIGFIGVVTTEIPNLVLKQNYEHYQFLDVAETIAKYAKELQEQHVHA
IVVLAHVPATSKDGVVDHEMATVMEKVNQIYPEHSIDIIFAGHNHQYTNGTIGKTRIVQALSQGKAYADVRGT
LDTDTNDFIKTPSANVVAVAPGIKTENSDIKAIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNL
ATTAQLTIAKKTFPTVDFAMTNNGGIRSDLVVKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDE
NQTYFLQMSGLTYTYTDNDPKNSDTPFKIVKVYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAI
NTDTEAFITYITNLEASGKTVNATIKGVKNYVTSNLESSTKVNSAGKHSIISKVFRNRDGNTVSSEVISDLLT
STENTNNSLGKKETTTNKNTISSSTLPIT

4.2 Antigenic fragment Spy0872-2

- > Spy0872-2 / SF370 (serotype 1); SEQ ID NO: 7
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLATTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIISKVFRNRDGNTVSSEVISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT
 - 4.3 Homologous sequences of other S. pyogenes isolates and/or serotypes
- > Spy0872-2 / Schmitz 1/7 (serotype 4); SEQ ID NO: 121
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWEAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIIIISKVFRNRDGNIVSSEIISDLLTSTENTNNSFGKKEITTNKNTISNSTLPIT
- > Spy0872-2 / Schmitz 1/39 (serotype 12); SEQ ID NO: 122
 40 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIISKVFRNRDGNIVSSEIISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT
- > Spy0872-2 / Schmitz 1/55 (serotype 118); SEQ ID NO: 123
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDIPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIISKVFRNRDGNIVSSEVISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT
- > Spy0872-2 / Schmitz 1/56 (serotype 28); SEQ ID NO: 124
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIISKVFRNRDGNIVSSEIISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT
- > Spy0872-2 / Schmitz 1/94 (serotype 1); SEQ ID NO: 125
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLATTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY

VTSNLESSTKVNSAGKHSIISKVFRNRDGNTVSSEVISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT

- > Spy0872-2 / Schmitz 1/253 (serotype 49); SEQ ID NO: 126
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTFTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIISKVFRNRDGNIVSSEIISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT
- > Spy0872-2 / Schmitz 1/176 (serotype 83); SEQ ID NO: 127

 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLVGAINTDTEAFITYITNLQASGKTVNATIKGVKNY
 VTSNLERSTKINSAGKHSIISKVFRNRDGNIVSSEVISDLLTSTENTNNSFGKKETTTNKNTISNSTLPIT
- > Spy0872-2 / Schmitz 1/177 (serotype 22); SEQ ID NO: 128
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWGAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIISKVFRNRDGNIVSSEIISDLLTSTENTNNSLGKKETTTNKNTISSSTLPIT
- > Spy0872-2 / Schmitz 1/22 (serotype 4); SEQ ID NO: 130
 AIINHANDIVKTVTERKIGTATNSSTISKTENIDKESPVGNLVTTAQLTIAKKTFPTVDFAMTNNGGIRSDLV
 VKNDRTITWEAAQAVQPFGNILQVIQMTGQHIYDVLNQQYDENQTYFLQMSGLTYTYTDNDPKNSDTPFKIVK
 VYKDNGEEINLTTTYTVVVNDFLYGGGDGFSAFKKAKLIGAINTDTEAFITYITNLEASGKTVNATIKGVKNY
 VTSNLESSTKVNSAGKHSIIIISKVFRNRDGNIVSSEIISDLLTSTENTNNSFGKKEITTNKNTISNSTLPIT

5. Further Sequences

- > Spy0488 / SF370 (serotype 1); SEQ ID NO: 8
 LRQIQSIRLIDVLELAFGVGYKEETTSQFSSDQPSQVVLYRGEANTVRFAYTNQMSLMKDIRIALDGSDKSLT
 AQIVPGMGHVYEGFQTSARGIFTMSGVPESTVPVANPNVQTKYIRYFKVIDDMHNTMYKGTVFLVQPQAWKYT
 MKSVDQLPVDDLNHIGVAGIERMTTLIKNAGALLTTGGSGAFPDNIKVSINPKGRQATITYGDGSTDIIPPAV
 LWKKGSVKEPTEADQSVGTPTPGIPGKFKRDQSLNEHEAMVNVEPLSHVVKDNIKVIDEKSTGRFEPFRPNED
 EKEKPASDVKVRPAEVGSWLEPATALPSVEMSAEDRLKS
- > Spy0895 / SF370 (serotype 1); SEQ ID NO: 9
 TNNQTLDILLDVYAYNHAFRIAKALPNIPKTALYLLEMLKERRELNLAFLAEHAAENRTIEDQYHCSLWLNQS
 LEDEQIANYILDLEVKVKNGAIIDFVRSVSPILYRLFLRLITSEIPNFKAYIFDTKNDQYDTWHFQAMLESDH
 EVFKAYLSQKQSRNVTTKSLADMLTLTSLPQEIKDLVFLLRHFEKAVRNPLAHLIKPFDEEELHRTTHFSSQA
 FLENIITLATFSGVIYRREPFYFDDMNAIIKKELSLWRQSIV
- > Spy1536 / SF370 (serotype 1); SEQ ID NO: 131
 IEMPGGAYDIRTVLQVNGKEDKRKGAYQFVAVGISRASLAQLLYAWLTPFTEISTAEDTTGGYSDADFLRINQ
 FYMETSQNAAIYQALSLAGKPVTLDYKGVYVLDVNNESTFKGTLHLADTVTGVNGKQFTSSAELIDYVSHLKL
 GDEVTVQFTSDNKPKKGVGRIIKLKNGKNGIGIALTDHTSVNSEDTVIFSTKGVGGPSAGLMFTLDIYDQITK
 EDLRKGRTIAGTGTIGKDGEVGDIGGAGLKVVAAAEAGADIFFVPNNPVDKEIKKVNPNAISNYEEAKRAAKR
 LKTKMKIVPVTTVQEALVYLRK
- > Spy1666 / SF370 (serotype 1); SEQ ID NO: 132
 TKEFHHVTVLLHETVDMLDIKPDGIYVDATLGGSGHSAYLLSKLGEEGHLYCFDQDQKAIDNAQVTLKSYIDK
 GQVTFIKDNFRHLKARLTALGVDEIDGILYDLGVSSPQLDERERGFSYKQDAPLDMRMDRQSLLTAYEVVNTY
 PFNDLVKIFFKYGEDKFSKQIARKIEQARAIKPIETTTELAELIKAAKPAKELKKKGHPAKQIFQAIRIEVND
 ELGAADESIQDAMELLALDGRISVITFHSLEDRLTKQLFKEASTVDVPKGLPLIPEDMKPKFELVSRKPILPS
 60 HSELTANKRAHSAKLRVAKKIRK

> Spy1727 / SF370 (serotype 1); SEQ ID NO: 10 VTTTEQELTLTPLRGKSGKAYKGTYPNGECVFIKLNTTPILPALAKEQIAPQLLWAKRMGNGDMMSAQEWLNG RTLTKEDMNSKQIIHILLRLHKSKKLVNQLLQLNYKIENPYDLLVDFEQNAPLQIQQNSYLQAIVKELKRSLP EFKSEVATIVHGDIKHSNWVITTSGMIFLVDWDSVRLTDRMYDVAYLLSHYIPRSRWSEWLSYYGYKNNDKVM QKIIWYGQFSHLTQILKCFDKRDMEHVNQEIYALRKFREIFRKK

SEQUENCE DATA FOR DNA SEQUENCES

1. Spy0269

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1.1 Full length Spy0269

> Spy0269 / SF370 (serotype 1); SEQ ID NO: 133 ATGGACTTAGAACAAACGAAGCCAAACCAAGTTAAGCAGAAAATTGCTTTAACCTCAACAATTGCTTTATTGA GTGCCAGTGTAGGCGTATCTCACCAAGTCAAAGCAGATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATAC 15 TCACGACGATAGTTTACCAAAACCAGAAACAATTCAAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACT CTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGCTACCGCTCTGACAAAAACTACTGCTGAAAATCAACCACT TAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTAACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAG TAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAACATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCAT AATGCTCAAGCAGATCAACATTCAAAAGAGACTGCATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTA 20 CTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACGTCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAG CAATCCTGATGCTATCACTAAAGCAGCTCAAACGGCTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAG AAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAAAGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGA AAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGTCGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCAT TGTGGGTAATAATACCATGAAAGCACCGCAAGGCTATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGT 25 TATATTGGATCAGCTAGTTACAATAATTATTACAAAGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAG GTAATCAATTAAATCAATACCAAGATATTCCAGCAGATCGTAATCGCTTTGTTGATCCCGATAATTTGACACC AGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTCACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCA CCAGTTACTGTTACAGCAGGATCACAAGAATTTGCAAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTA ATACAAGACCATCATTTGTCTACGGACAGCCAGGGGTATCAGGGCATTATGGTGTTGGGCCTCATGATAAAAC 30 TATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCATTCGAAATGATGATAACATGTACGAGAATATCGGTGCT TTTAACGATGTGCATACTGTGAATGGTATTAAACGTGGTATTTATGACAGTATCAAGTATATGCTCTTTACAG ATCATTTACACGGAAATACATACGGCCATGCTATTAACTTTTTACGTGTAGATAAACATAACCCTAATGCGCC ATTGCTAACCATCAACGCTTTAATAAGACCCCTATAAAAGCCGTTGGAAGTACAAAAGATTATGCCCAAAGAG 35 TAGGCACTGTATCTGATACTATTGCAGCGATCAAAGGAAAAGTAAGCTCATTAGAAAAATCGTTTGTCGGCTAT AAGCAGTCAGACAGCTTAAATCTCCAAGTGAGACAATTAAATGATACTAAAGGTTCTTTGAGAACAGAATTAC AGCCGCACTGCACCAGACAGAAGCCTTAGCAGAGCAAGCCGCAGCCAGAGTGACAGCACTGGTGGCTAAAAAA 40 GCTCATTTGCAATATCTAAGGGACTTTAAATTGAATCCTAACCGCCTTCAAGTGATACGTGAGCGCATTGATA ATACTAAGCAAGATTTGGCTAAAACTACCTCATCTTTGTTAAATGCACAAGAAGCTTTAGCAGCCTTACAAGC TAAACAAAGCAGTCTAGAAGCTACTATTGCTACCACAGAACACCAGTTGACTTTGCTTAAAAACCTTAGCTAAC GAAAAGGAATATCGCCACTTAGACGAAGATATAGCTACTGTGCCTGATTTGCAAGTAGCTCCACCTCTTACGG GCGTAAAACCGCTATCATATAGTAAGATAGATACTACTCCGCTTGTTCAAGAAATGGTTAAAGAAACGAAACA 45 ACTATTAGAAGCTTCAGCAAGATTAGCTGCTGAAAATACAAGTCTTGTAGCAGAAGCGCTTGTTGGCCAAACC TCTGAAATGGTAGCAAGTAATGCCATTGTGTCTAAAATCACATCTTCGATTACTCAGCCCTCATCTAAGACAT CTTATGGCTCAGGATCTTCTACAACGAGCAATCTCATTTCTGATGTTGATGAAAGTACTCAAAGAGCTCTTAA AGCAGGAGTCGTCATGTTGGCAGCTGTCGGCCTCACAGGATTTAGGTTCCGTAAGGAATCTAAGTGA

1.2 Antigenic fragment Spy0269-1

> Spy0269-1 / SF370 (serotype 1); SEQ ID NO: 11
GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC
AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC
TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA
ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC
ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC
ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG

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1.3 Homologous sequences of other S. pyogenes isolates and/or serotypes

> Spy0269-1 / Schmitz 2/14 (serotype 1); SEQ ID NO: 134 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTGAAATCAACAACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA 20 ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA 25 AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATTCCAGCA GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC 30 ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTTACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG 35 AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT

> Spy0269-1 / Schmitz 1/156 (serotype 4); SEQ ID NO: 135 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC 40 TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG 45 CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATTCCAGCA 50 GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGGAGACAATTAGGTCTACCACCAGTTACTGTCACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG 55 TGGTATTTATGACAGTATCAAGTATATGCTCTTTACAGATCATTTACACGGAAATACATATGGTCATGCTATT AACTTTTTACGTGTAGATAAACATAACCCTAAGGCGCCTGTT

> Spy0269-1 / Schmitz 1/59 (serotype 12); SEQ ID NO: 136
GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC

AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAAAAGCTTGC TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG 5 TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA 10 AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATTCCAGCA GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTTACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA 15 TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT

- > Spy0269-1 / Schmitz 1/177 (serotype 22); SEQ ID NO: 137 20 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGAAGCAGTTGAAAAAGCTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTAAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC 25 ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGTAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT 30 ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATTCCAGCA GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTCACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG 35 GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG TGGTATTTATGACAGTATCAAGTATATGCTCTTTACAGATCATTTACACGGAAATACATATGGCCATGCTATT AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT
- 40 > Spy0269-1 / Schmitz 1/43 (serotype 22); SEQ ID NO: 138 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGAAGCAGTTGAAAAAGCTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTAAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC 45 ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGTAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAAGAGGCAGAACTTAGT 50 CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATTCCAGCA GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTCACAGCAGGATCACAAGAATTTGC 55 AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG TGGTATTTATGACAGTATCAAGTATATGCTCTTTACAGATCATTTACACGGAAATACATATGGCCATGCTATT 60 AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT

- > Spy0269-1 / Schmitz 1/136 (serotype 25); SEQ ID NO: 139 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA 5 ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA 10 AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATTCCAGCA GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC 15 ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTTACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG 20 AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT
- > Spy0269-1 / Schmitz 1/85 (serotype 28); SEQ ID NO: 140 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC 25 TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG 30 CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGatattccagca 35 gatcgtaatcgctttGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTTACAGCAGGATCACaagaatttgc aagattacttagtaccagctataagaaaactcatggtaatacaagaccatcatttgtctACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG 40 AACTTTTTACGTGTAGATAAACATAACCCTAATGCGCCTGTT
- > Spy0269-1 / Schmitz 2/50 (serotype 28); SEQ ID NO: 141 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC 45 AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGTAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG 50 TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA 55 AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATtCCAGCA GATCGTAATCGCTTTGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTTACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAGACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA 60

TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG TGGTATTTATGACAGTATCAAGTATAŢGCTCTTTACAĢATCATTTACACGGAAATACATACGGCCATGCTATT AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT

- > Spy0269-1 / Schmitz 1/123 (serotype 49); SEQ ID NO: 142 5 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGATGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAATGAACAAAAAGCTTTA ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC 10 ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGTAATAATACCATGAAAGCACCGCAAGGCT 15 ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGATATCCAGCA GAtcqtaatcqctttGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGGAGACAATTAGGTCTACCACCAGTTACTGTTACAGCAGGATCACAAGAATTTGC AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAACCAGGG 20 GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG TGGTATTTATGACAGTATCAAGTATATGCTCTTTACAGATCATTTACACGGAAATACATATGGCCATGCTATT AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT
- 25 > Spy0269-1 / Schmitz 1/176 (serotype 83); SEQ ID NO: 143 GATGATAGAGCCTCAGGAGAAACGAAGGCGAGTAATACTCACGACGATAGTTTACCAAAACCAGAAACAATTC AAGAGGCAAAGGCAACTATTGAAGCAGTTGAAAAAACTCTCAGTCAACAAAAAGCAGAACTGACAGAGCTTGC TACCGCTCTGACAAAAACTACTGCTGAAATCAACCACTTAAAAGAGCAGCAAGATAACGAACAAAAAGCTTTA 30 ACCTCTGCACAAGAAATTTACACTAATACTCTTGCAAGTAGTGAGGAGACGCTATTAGCCCAAGGAGCCGAAC ATCAAAGAGAGTTAACAGCTACTGAAACAGAGCTTCATAATGCTCAAGCAGATCAACATTCAAAAGAGACTGC ATTGTCAGAACAAAAAGCTAGCATTTCAGCAGAAACTACTCGAGCTCAAGATTTAGTGGAACAAGTCAAAACG TCTGAACAAAATATTGCTAAGCTCAATGCTATGATTAGCAATCCTGATGCTATCACTAAAGCAGCTCAAACGG CTAATGATAATACAAAAGCATTAAGCTCAGAATTGGAGAAGGCTAAAGCTGACTTAGAAAATCAAAAAGCTAA 35 AGTTAAAAAGCAATTGACTGAAGAGTTGGCAGCTCAGAAAGCTGCTCTAGCAGAAAAAGAGGCAGAACTTAGT CGTCTTAAATCCTCAGCTCCGTCTACTCAAGATAGCATTGTGGGGTAATAATACCATGAAAGCACCGCAAGGCT ATCCTCTTGAAGAACTTAAAAAATTAGAAGCTAGTGGTTATATTGGATCAGCTAGTTACAATAATTATTACAA AGAGCATGCAGATCAAATTATTGCCAAAGCTAGTCCAGGTAATCAATTAAATCAATACCAAGatATTCCAGCA GatcgtaatcgctttGTTGATCCCGATAATTTGACACCAGAAGTGCAAAATGAGCTAGCGCAGTTTGCAGCTC ACATGATTAATAGTGTAAGAAGACAATTAGGTCTACCACCAGTTACTGTCACAGCAGGATCACAAGAATTTGC 40 AAGATTACTTAGTACCAGCTATAAGAAAACTCATGGTAATACAAGACCATCATTTGTCTACGGACAGCCAGGG GTATCAGGGCATTATGGTGTTTGGGCCTCATGATAAAACTATTATTGAAGACTCTGCCGGAGCGTCAGGGCTCA TTCGAAATGATGATAACATGTACGAGAATATCGGTGCTTTTAACGATGTGCATACTGTGAATGGTATTAAACG TGGTATTTATGACAGTATCAAGTATATGCTCTTTACAGATCATTTACACGGAAATACATATGGCCATGCTATT 45 AACTTTTTACGTGTAGATAAACGTAACCCTAATGCGCCTGTT

2. Spy0292

2.1 Full length Spy0292

10 2.2 Antigenic fragment Spy0292-1

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2.3 Homologous sequences of other S. pyogenes isolates and/or serotypes

- > Spy0292-1 / Schmitz 1/76 (serotype 22); SEQ ID NO: 149
 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG

2.4 Antigenic fragment Spy0292-3

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> Spy0292-3 / SF370 (serotype 1); SEQ ID NO: 13 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG ATGCTAAAGAAGTTGTCCCAGTCGCCTCAGTCAGTAAGCTCTTGACAACCTATCTGGTTTACAAAGAAGTTTC TAAGGGCAAGCTAAATTGGGATAGTCCTGTAACTATTTCTAACTACCCTTATGAACTCACTACAAACTATACT ATTAGTAACGTTCCTCTTGATAAGAGAAAAATATACCGTTAAAGAACTTTTAAGTGCGTTAGTTGTTAATAACG CCAATAGCCCCGCTATTGCTTTAGCTGAAAAAAATAGGCGGAACCGAACCCAAATTTGTTGACAAAATGAAAAA

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2.5 Homologous sequences of other S. pyogenes isolates and/or serotypes

- 10 > Spy0292-3 / Schmitz 1/39 (serotype 12); SEQ ID NO: 155 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG ATGCTAAAGAAGTTGTCCCAGTCGCCTCAGTCAGTAAGCTCTTGACAACCTATCTGGTTTACAAAGAAGTTTC TAAGGCCAAGCTAAATTGGGATAGTCCTGTAACTATTTCTAACTACCCTTATGAACTCACTACAAACTATACT ATTAGTAACGTTCCTCTTGATAAGAGAAAATATACCGTTAAAGAACTTTTAAGTGCGTTAGTTGTTAATAACG 15 CCAATAGCCCCGCTATTGCTTTAGCTGAAAAAATAGGCGGAACCGAACCCAAATTTGTTGACAAAATGAAAAA GCTAATACTTATCCTAATACAGAACcagaTGATGAAAATTGTTTTTTGCGCCACTGATTTAGCTATTATTGCCA GGCATCTCTTATTAGAATTTCCAGAAGTACTGAAATTATCTAGCAAATCCTCCACTATTTTTGATGGACAAAC CATTTACAGTTATAATTACATGCTTAAAGGCATGCCTTGTTATCGAGAAGGCGTGGATGGTCTTTTTGTTGGT 20 TATTCTAAAAAAGCCGGTGCTTCTTTTGTAGCTACTAGTGTCGAAAATCAAATGAGGGTTATTACAGTAGTTT AATTAATTTTCAAAAAGTCCAGTTAATTGAA
- > Spy0292-3 / Schmitz 1/55 (serotype 118); SEQ ID NO: 156 25 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG ATGCTAAAGAAGTTGTCCCAGTCGCCTCAGTCAGTAAGCTCTTGACAACCTATCTGGTTTACAAAGAAGTTTC TAAGGGCAAGCTAAATTGGGATAGTCCTGTAACTATTTCTAACTACCCTTATGAACTTACTACAAACTATACT ATTAGTAACGTTCCTCTTGATAAGAGAAAATATACCGTTAAAGAACTTTTAAGTGCGTTAGTTGTTAATAACG CCAATAGCCCCGCTATTGCTTTAGCTGAAAAAATAGGCGGAACCGAACCCAAATTTGTTGACAAAATGAAAAA 30 GCTAATACTTATCCTAATACAGAACCAGATGATGAAAATTGTTTTTTGCGCCACTGATTTAGCTATTATTGCCA GGCATCTCTTATTAGAATTTCCAGAAGTACTGAAATTATCTAGCAAATCCTCCACTATTTTTTGATGGACAAAC CATTTACAGTTATAATTACATGCTTAAAGGCATGCCTTGTTATCGAGAAGGCGTGGATGGTCTCTTTGTCGGT 35 TATTCTAAAAAAGCCGGTGCTTCTTTTGTAGCTACTAGTGTCGAAAATCAAATGAGGGTTATTACAGTAGTTT AATTAATTTTCAAAAAGTCCAGTTAATTGAA
- > Spy0292-3 / Schmitz 1/56 (serotype 28); SEQ ID NO: 157 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG 40 ATACTAAAGAAGTTGTCCCAGTCGCCTCAGTCAGTAAGCTCTTGACAACCTATCTGGTTTACAAAGAAGTTTC TAAGGGCAAGCTAAATTGGGATAGTCCTGTAACTATTTCTAACTACCCTTATGAACTCACTACAAACTATACT ATTAGTAACGTTCCTCTTGATAAGAGAAAATATACCGTTAAAGAACTTTTAAGTGCGTTAGTTGTTAATAACG CCAATAGCCCCGCTATTGCTTTAGCTGAAAAAATAGGCGGAACCGAACCCAAATTTGTTGACAAAATGAAAAA 45 GCTAATACTTATCCTAATACAGAACCAGATGATGAAAATTGTTTTTTGCGCCACTGATTTAGCTATTATTGCCA GGCATCTCTTATTAGAATTTCCAGAAGTACTGAAATTATCTAGCAAATCCTCCACTATTTTTGATGGACAAAC CATTTACAGTTATAATTACATGCTTAAAGGCATGCCTTGTTATCGAGAAGGCGTGGATGGTCTTTTTGTTGGT TATTCTAAAAAAGCCGGTGCTTCTTTTGTAGCTACTAGTGTCGAAAATCAAATGAGGGTTATTACAGTAGTTT 50 AATTAATTTTCAAAAAGTCCAGTTAATTGAA

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- > Spy0292-3 / Schmitz 1/94 (serotype 1); SEQ ID NO: 161 35 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG ATGCTAAAGAAGTTGTCCCAGTCGCCTCAGTCAGTAAGCTCTTGACAACCTATCTGGTTTACAAAGAAGTTTC TAAGGGCAAGCTAAATTGGGATAGTCCTGTAACTATTTCTAACTACCCTTATGAACTCACTACAAACTATACT ATTAGTAACGTTCCTCTTGATAAGAGAAAATATACCGTTAAAGAACTTTTAAGTGCGTTAGTTGTTAATAACG CCAATAGCCCCGCTATTGCTTTAGCTGAAAAAATAGGCGGAACCGAACCCAAATTTGTTGACAAAATGAAAAA 40 GCTAATACTTATCCTAATACAGAACCAGATGATGAAAATTGTTTTTTGCGCCACTGATTTAGCTATTATTGCCA GGCATCTCTTATTAGAATTTCCAGAAGTACTGAAATTATCTAGCAAATCCTCCACTATTTTTTGCTGGACAAAC CATTTACAGTTATAATTACATGCTTAAAGGCATGCCTTGTTATCGAGAAGGCGTGGATGGTCTTTTTTGTTGGT TATTCTAAAAAAGCCGGTGCTTCTTTTGTAGCTACTAGTGTCGAAAATCAAATGAGGGTTATTACAGTAGTTT 45 AATTAATTTTCAAAAAGTCCAGTTAATTGAA

AATTAATTTTCAAAAAGTCCAGTTAATTGAA

> Spy0292-3 / Schmitz 1/144 (serotype 76); SEQ ID NO: 163 GAAGAGTATTCGGTAACTGCTAAGCATGCGATTGCCGTTGACCTTGAAAGTGGCAAAGTTTTATACGAAAAAG ATGCTAAAGAAGTTGTCCCTGTCGCCTCAGTCAGTAAGCTCTTGACAACCTATCTGGTTTACAAAGAAGTTTC 5 TAAGGGCAAGCTAAATTGGGATAGTCCTGTAACTATTTCTAACTACCCTTATGAACTCACTACAAACTATACT ATTAGTAACGTTCCTCTTGATAAGAGAAAATATACCGTTAAAGAACTTTTAAGTGCGTTAGTTGTTAATAACG CCAATAGCCCCGCTATTGCTTTAGCTGAAAAAATAGGCGGAACCGAACCCAAATTTGTTGACAAAATGAAAAA GCTAATACTTATCCTAATACAGAaccagaTGATGAAAATTGTTTTTTGCGCCACTGATTTAGCTATTATTGCCA 10 GGCATCTCTTATTAGAATTTCCAGAAGTACTGAAATTATCTAGCAAATCCTCCACTATTTTTTGATGGACAAAC CATTTACAGTTATAATTACATGCTTAAAGGCATGCCTTGTTATCGAGAAGGCGTGGATGGTCTTTTTTGTTGGT TATTCTAAAAAAGCCGGTGCTTCTTTTGTAGCTACTAGTGTCGAAAATCAAATGAGGGTTATTACAGTAGTTA AATTAATTTTCAAAAAGTCCAGTTAATTGAA 15

3. Spy0416A

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3.1 Full length Spy0416A

> Spy0416A / SF370 (serotype 1); SEQ ID NO: 165 35 GCAGATGAGCTAAGCACAATGAGCGAACCAACAATCACGAATCACGCTCAACAACAAGCGCAACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCAAAATCTCAAGACACATCACAAATCACTCTCAAGACAAATCGTGAAAA AATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA 40 TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG 45 AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACATCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCGACAAATCCAGAC 50 TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA TTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTATTCAGA GTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAG TCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCTCTGGGAGTACTTATTTTTAATAACAAGCCTGG 55 TCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTT GGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCAC CGAGTCAAAAAGGCAATGAAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGA CATTACTGCACCAGGTGGCGATATCTATTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGT ATGGCCTCTCCTCAGATTGCTGGCGCCAGCCTTTTGGTCAAACAATACCTAGAAAAAGACTCAGCCAAACTTGC 60

CAAAAGAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGCAATGCTCAAATTCATGTTAATCCAGAGAC
AAAAACGACCACCTCACCGCGTCAGCAAGGGGCAGGATTACTTAATATTGACGGAGCTGTCACTAGCGGCCTT
TATGTGACAGGAAAAGACAACTATGGCAGTATATCATTAGGCAACATCACAGATACGATGACGTTTGATGTGA
CTGTTCACAACCTAAGCAATAAAGACAAAACATTACGTTATGACACAGAATTGCTAACAGATCATGTAGACCC
ACAAAAGGGCCGCTTCACTTTGACTTCTCACTCCTTAAAAACGTACCAAGGAGGAGAAGTTACAGTCCCAGCC
AATGGAAAAGTGACTGTAAGGGTTACCATGGATGTCTCACAGTTCACAAAAGAGCTAACAAAACAGATGCCAA
ATGGTTACTATCTAGAAGGTTTTGTCCGCTTTAGAGATAGTCAAGATGACCAACTAAATAGAGTAAACATTCC
TTTTGTTGGTTTTTAAAGGGCAATTTGAAAACTTAGCAGTTGCAGAAGAGTCCATTTACAGATTAAAATCTCAA
GGCAAAACTGGTTTTTACTTTGATGAATCAGGTCCAAAAGACCGATATCTATGTCGGTAAACACTTTACAGGAC

3.2 Antigenic fragment Spy0416A-1

AAAAACGACCACCTCACCGCGTCAGCAAGGGGCA

TTGTCACTCTTGGTTCAGAG

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> Spy0416A-1 / SF370 (serotype 1); SEQ ID NO: 14 GCAGATGAGCTAAGCACAATGAGCGAACCAACAATCACGAATCACGCTCAACAACAAGCGCAACATCTCACCA 15 ATACAGAGTTGAGCTCAGCTGAATCAAAATCTCAAGACACATCACAAATCACTCTCAAGACAAATCGTGAAAA AATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA 20 AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT 25 TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACATCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCGACAAATCCAGAC TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA 30 TTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTATTCAGA GTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAG TCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCTCTGGGAGTACTTATTTTTAATAACAAGCCTGG TCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTT 35 GGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCAC CGAGTCAAAAAGGCAATGAAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGA CATTACTGCACCAGGTGGCGATATCTATTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGT ATGGCCTCTCCTCAGATTGCTGGCGCCAGCCTTTTGGTCAAACAATACCTAGAAAAAGACTCAGCCAAACTTGC CAAAAGAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGGCAATGCTCAAATTCATGTTAATCCAGAGAC 40

3.3 Homologous sequences of other S. pyogenes isolates and/or serotypes

> Spy0416A-1 / Schmitz 1/7 (serotype 4); SEQ ID NO: 166 45 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACGCTCAACAACAACAACGACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCACAATCCCCAGACACATCACAAATCACTCCCAAGACAAATCGTGAAAA AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA 50 TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGGGGATTTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC AAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACGAGTCAC 55 ACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTAGG AATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTC TTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATG GGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATCAGTTGT TGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGACTATGGT 60

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> Spy0416A-1 / Schmitz 1/39 (serotype 12); SEQ ID NO: 167 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACACTCAACAACAAGCGCAACATCTCACCA 15 ATACAGAGTTGAGCTCAGCTGAATCAAAACCTCAAGACACATCACAAATCACTCTCAAGACAAATCGTGAAAA AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA 20 AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT 25 TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGAC TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA 30 TTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTaTTCAGA GTCTGTCGActttaaaGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCaAAGAG TCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGAGTACTTATTTTTAATAACAAGCCTGG TCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTT 35 GGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCAC CGAGTCAAAAAGGCAATGAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGA CATTACTGCACCAGGTGGCGATATCTACTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGT ATGGCCTCTCCTCAGATTGCTGGCGCCAGCCTTTTGGTCAAACAATACCTAGAAAAAGACTCAGCCAAACTTGC CAAAAGAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGGCAATGCTCAAATTCATGTTAATCCAGAGAC 40 AAAAACGACCACCTCACCGCGTCAGCAAGGGGCA

> Spy0416A-1 / Schmitz 1/55 (serotype 118); SEQ ID NO: 168 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACGCTCAACAACAAGCGCCACCTCTCACCA ATACAGAGTTGAGCTCAGCTGAATCACAACCTCAAGACACATCACAAGTAACTCCAGAGACAAATCGTGAAAA 45 AATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA 50 AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAA 55 TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCCTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCGACAAATCCAGAC TATGGTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTAGCAGCTATAAACAGTAAGTGGGTGA TTCAACGTCTAATGACGGTCAAAGAATTGGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTaTTCAGA 60

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> Spy0416A-1 / Schmitz 1/56 (serotype 28); SEQ ID NO: 169 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACGCTCAACAACAAGCGCAACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCACAATCCCCAGACACATCACAAATCACTCCCAAGATAAATCGTGAAAA 15 AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGTACAAAGGACAAGGTAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC 20 AAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACGAGTCAC ACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGTAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTAGG AATTGCACCAGAGACCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTC TTTATCAAAGCTATCGAAGATGCCGTGGCCTTAGGAGCAGATGTGATCAACCTGAGTCTTGGGACCGCTAATG 25 GTGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATCAGTTGT TGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGACTATGGT TTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAAC GTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTaTTCAGAGTCTGT CGACTTLAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAGTCAACT 30 GATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAACCTATG ACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGAGTACTTATTTTTAATAACAAGCCTGGTCAATC AAACCGCTCAATGCGCCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTTGGTAAG GCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCACCGAGTC AAAAAGGCAATGAAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGACATTAC 35 TGCACCAGGGGGTGATATCTACTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGTATGGCC TCTCCTCAGATTGCTGGCGCCAGCCTTTTGGTCAAACAATACCTAGAAAAGACTCAGCCAAAACTTGCCAAAAG AAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGCAATGCTCAAATTCATGTTAATCCAGAGACAAAAAC GACCACCTCACCGCGTCAGCAAGGGGCA

> Spy0416A-1 / Schmitz 1/94 (serotype 1); SEQ ID NO: 170 GCAGATGAGCTAAGCACAATGAGCGAACCAACAATCACGAATCACGCTCAACAACAAGCGCAACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCAAAATCTCAAGACACATCACAAATCACTCTCAAGACAAATCGTGAAAA AATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA 45 AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC 50 GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACATCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATC 55 AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCGACAAATCCAGAC TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA TTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTATTCAGA GTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCaAAGAG TCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA 60

- > Spy0416A-1 / Schmitz 1/253 (serotype 49); SEQ ID NO: 171 10 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACGCTCAACAACAACAAGCGCAACCTCTCACCA ATACAGAGTTGAGCTCAGCTGAATCACAATCCCCAGACATATCACAAGTAACTCCAGAGACAAATCGTGAAAA AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA 15 TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG 20 AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGAC 25 TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA TTCAACGTCTAATGACGGTCAAAGGATTAGAAAACCGTGCCGATTTAaACCATGGTAAAGCCATCTATTCAGA GTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAG TCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGACTACTTATTTTTAATAACAAGTCTGG 30 TCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTT GGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCAC CGAGTCAAAAAGGCAATGAAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGA CATTACTGCACCAGGTGGCGATATCTACTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGT ATGCCTCTCCTCAGATTGCTGGCGCCCAGCCTTTTGGTCAAACAATACCTAGAAAAAGACCCAGCCAAACTTGC 35 CAAAAGAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGGCAATGCTCAAATTCATGTTAATCCAGAGAC AAAAACAACCACCTCACCGCGTCAGCAAGGGGCA
- > Spy0416A-1 / Schmitz 1/174 (serotype 6); SEQ ID NO: 172 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACGCTCAACAACAACAACCACCACCA 40 ATACAGAGTTGAGCTCAGCTGAATCAAAACCTCAAGACACATCACAAATCACTCCCAAGACAAATCGTGAAAA AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA 45 AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT 50 TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGAC TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA 55 TTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTaTTCAGA GTCTGTCGACTTTAAAAACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAG TCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGAGTACTTATTTTTAATAACAAACCTGG TCAATCAAACCGCTCAATGCGCCTAACATCTAATGGGATGGGAATACCATCTGCTTTCATATCGCACGAATTT 60

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> Spy0416A-1 / Schmitz 1/176 (serotype 83); SEQ ID NO: 173 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACACTCAACAACAAGCGCAACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCAAAACCTCAAGACACATCACAAATCACTCTCAAGACAAATCGTGAAAA 10 AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA 15 AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAA 20 TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGAC TATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA TTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTATTCAGA 25 GTCTGTCGACTTTAAAAACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAG TCAACTGATGCGGGTTATAAAGCACAAGACGTTAAAGGTAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGAGTACTTATTTTTAATAACAAGCCTGG TCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTT GGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCAC 30 CGAGTCAAAAAGGCAATGAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGA CATTACTGCACCAGGTGGCGATATCTACTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGT ATGGCCTCTCCTCAGATTGCTGGCGCCCAGCCTTTTGGTCAAACAATACCTAGAAAAAGACTCAGCCAAACTTGC CAAAAGAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGGCAATGCTCAAATTCATGTTAATCCAGAGAC AAAAACGACCACCTCACCGCGTCAGCAAGGGGCA 35

> Spy0416A-1 / Schmitz 1/234 (serotype 44); SEQ ID NO: 174 GCAGATGAGCTAAGCACAATGAGTGAACCAACAATCACGAATCACGCTCAACAACAAGCGCAACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCAAAATCTCAAGACACATCACAAATCACTCCCAAGACAAATCGTGAAAA 40 AATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC 45 GTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACG AGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTT TTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAA TCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCG 50 CTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATC AGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGACCCATTGGCAACAAATCCAGAC TATGGTTTGGTTGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGA TTCAACGTCTAATGACGGTCAAAGAATTGGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTaTTCAGA GTCTGTCGACTTtAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAG 55 TCAACTGATGCGGGTTATAAAGCACAAGACGTTAAAGATAAAATTGCTTTAATTGAACGTGATCCCAATAAAA CCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGGGGTACTTATTTTTAATAACAAGCCTGG TCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTT GGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCAC CGAGTCAAAAAGGCAATGAAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGA 60

CATTACTGCACCAGGCGGCGATATCTACTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGT ATGGCCTCTCCTCAGATTGCTGGCGCCCAGCCTTTTGGTCAAACAATATCTAGAAAAAAGACTCAGCCAAACTTGC CAAAAGAAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGCAATGCTCAAATTCATGTTAATCCAGAGAC AAAAACGACCACCTCACCGCGTCAGCAAGGGGCA

5 > Spy0416A-1 / Schmitz 1/22 (serotype 4); SEQ ID NO: 175 GCAGATGAGCTAACCACAACGAGTGAACCAACAATCACGAATCACGCTCAACAACAAGCGCAACATCTCACCA ATACAGAGTTGAGCTCAGCTGAATCACAATCCCCAGACACATCACAAATCACTCCCAAGACAAATCGTGAAAA AATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACCGCCAGTCAATACAGATGTTCACGATTGGGTAA 10 AAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGCAAGGTTGTCGCAGTTATTGACACAGGGATCGA TCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAAAATCAAAAGAAGAAGACATGCTAGCA AAAATAGCGATAATATCAAAGAAAATCAATTCGGGGGATTTTGATGAGGACTGGGAAAACTTTGAGTTTGATGC 15 AAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGACGATGACACCAAATACGAGTCAC ACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTAGG AATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTC TTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATG GGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCTAAAAAAGCCGGTGTATCAGTTGT 20 TGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCATTGGCAACAAATCCAGACTATGGT TTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAAC GTCTAATGACGGCCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGTAAAGCCATCTATTCAGAGTCTGT CGACTTTAAAGACATAAAAGATAGCCTAGGTTATGATAAATCGCATCAATTTGCTTATGTCAAAGAGTCAACT GATGCGGGTTATAAAGCACAAGACGTTAAAGATAAAATTGCTTTAATTGAACGTGATCCCAATAAAACCTATG 25 ACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGAGTACTTATTTTTAATAACAAGCCTGGTCAATC AAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATACCATCTGCTTTCATATCGCACGAATTTGGTAAG GCCATGTCCCAATTAAATGGCAATGGTACAGGAAGTTTAGAGTTTGACAGTGTGGTCTCAAAAGCACCGAGTC AAAAAGGCAATGAAATGAATCATTTTTCAAATTGGGGCCTAACTTCTGATGGCTATTTAAAAACCTGACATTAC TGCACCAGGTGGCGATATCTACTCTACCTATAACGATAACCACTATGGTAGCCAAACAGGAACAAGTATGGCC 30 TCTCCTCAGATTGCTGGCGCCAGCCTTTTGGTCAAACAATACCTAGAAAAAGACTCAGCCAAAACTTGCCAAAAG AAAAAATTGCTGATATCGTTAAGAACCTATTGATGAGCAATGCTCAAATTCATGTTAATCCAGAGACAAAAAC GACCACCTCACCGCGTCAGCAAGGGGCA

35 3.4 Antigenic fragment Spy0416A-6

> Spy0416A-6 / SF370 (serotype 1); SEQ ID NO: 15 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC 40 TGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCATCAAAAAACACAAGATCTATCGTCCCCAAT CAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACA AACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAA GCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCA ACGACATCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGT 45 GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA AAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG ATCCATTGGCGACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTGGC AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAAC CATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTA 50

3.5 Homologous sequences of other S. pyogenes isolates and/or serotypes

GCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACG
TCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAA
CCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAAGCT
AAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCAT
TGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTAT
AAACAGTAAGTGGGTGATTCAACGTCTAATGACGGCCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGT
AAAGCCATCTATTCAGAGTCTGTCGActttaaagacataaaagatagccta

- > Spy0416A-6 / Schmitz 1/39 (serotype 12); SEQ ID NO: 177 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA 10 TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC TGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAGCCATCAAAAAACACAAGATCTATCGTCCCCAAT CAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACA AACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAA 15 GCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTGCCA ACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGT GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA AAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG ATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGC 20 AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAAC CATGGTAAAGCCATCTaTTCAGAGTCTGTCGActttaaaGACATAAAAGATAGCCTA
- > Spy0416A-6 / Schmitz 1/55 (serotype 118); SEQ ID NO: 178 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA 25 TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC TGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAGCCATCAAAAAACACAAGATCTATCGTCCCCAAT CAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACA AACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAA 30 GCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCA ACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCCTTAGGAGCAGATGT GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA AAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG ATCCATTGGCGACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTAGC 35 AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTGGAAAAACCGTGCCGATTTAAAC CATGGTAAAGCCATCTaTTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTA
- > Spy0416A-6 / Schmitz 1/56 (serotype 28); SEQ ID NO: 179 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA 40 TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC TGGGAAAACTTTGAGTTTGATGCAGAGCCAAAAGCCATCAAAAAACACAAGATCTATCGTCCCCAATCAACCC AGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGA CGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGTAAAGAAGCCGCT 45 GCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGACCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACG TCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCCTTAGGAGCAGATGTGATCAA CCTGAGTCTTGGGACCGCTAATGGTGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAGCT AAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCAT TGGCAACAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTGGCAGCTAT 50 AAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGT AAAGCCATCTaTTCAGAGTCTGTCGACTTtAAAGACATAAAAGATAGCCTA

GCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCA
ACGACATCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGT
GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA
AAAGCTAAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG
ATCCATTGGCGACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTGGC
AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAAACCGTGCCGATTTAAAC
CATGGTAAAGCCATCTATTCAGAGTCTGTCGACTttAAAAGACATAAAAGATAGCCTA

- > Spy0416A-6 / Schmitz 1/253 (serotype 49); SEQ ID NO: 181 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA 10 TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC TGGGAAAACTTTGAGTTTGATGCAGATGCAGAGCCAAAAGCCATCAAAAAACACAAGATCTATCGTCCCCAAT CAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACA AACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAA 15 GCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCA ACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGT GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA AAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG 20 ATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTGGC AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGGATTAGAAAACCGTGCCGATTTAaAC CATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTA
- > Spy0416A-6 / Schmitz 1/174 (serotype 6); SEQ ID NO: 182 25 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC TGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAGCCCATCAAAAAACACAAGATCTATCGTCCCCAAT CAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACA 30 AACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAA GCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCA ACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGT GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA AAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG 35 ATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTGGC AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAAACCGTGCCGATTTAAAC CATGGTAAAGCCATCTaTTCAGAGTCTGTCGACTTTAAAAAACATAAAAGATAGCCTA
- > Spy0416A-6 / Schmitz 1/176 (serotype 83); SEQ ID NO: 183 40 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTTGATGAGGAC TGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAGCCCATCAAAAAACACAAGATCTATCGTCCCCAAT CAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACA 45 AACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAA GCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCA ACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGT GATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAA AAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATG 50 ATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGTGGC AGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAAACCGTGCCGATTTAAAC CATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAAACATAAAAGATAGCCTA

> Spy0416A-6 / Schmitz 1/22 (serotype 4); SEQ ID NO: 185 GCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTGCTAAAGTAA 10 TGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGGGGATTTTGATGAGGAC AGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGACACAAACAGA CGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAAGAAGCCGCT 15 GCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTTGCCAACGACG TCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGATGTGATCAA CCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATTGAAAAAGCT AAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATGATGATCCAT TGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAACATCAGTGGCAGCTAT 20 AAACAGTAAGTGGGTGATTCAACGTCTAATGACGGCCAAAGAATTAGAAAACCGTGCCGATTTAAACCATGGT AAAGCCATCTATTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTA

3.6 Antigenic fragment Spy0416A-7

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> Spy0416A-7 / SF370 (serotype 1); SEQ ID NO: 16 TCACAAATCACTCTCAAGACAAATCGTGAAAAAGAGCAATCACAAGATCTAGTCTCTGAGCCAACCACAACTG AGCTAGCTGACACAGATGCAGCATCAATGGCTAATACAGGTTCTGATGCGACTCAAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG 30 TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCATCAAAAAACACAAGATCTATC GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT 35 AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACATCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA GCAATTGAAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGATCCATTGGCGACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAAC 40 ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCAAAGAGTCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAA AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCT 45 GGATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG \mathbf{T}

3.7 Homologous sequences of other S. pyogenes isolates and/or serotypes

GAAGCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTG
CCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGA
TGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATT
GAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATG
ATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGT
GGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGCCAAAGAATTAGAAAACCGTGCCGATTTA
AACCATGGTAAAGCCATCTATTCAGAGTCTGTCGACtttaaagacataaaagatagcctaggttatgataaAT
CGCATCAATTTGCTTATGTCAAaGAGTCAACTGATGCGGGTTATAAAGCACAAGACGTTAAAGATAAAATTGC
TTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTTGGCTAAGAAACATGGAGCCCTGGGA
GTACTTATTTTTAATAACAAGCCTGGTCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGCGAATAC
CATCTGCTTTCATATCGCACGAATTTGGTAAGGCCCATGTCCCCAATTAAATGGCAATGGTACAGGAAGT

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> Spy0416A-7 / Schmitz 1/39 (serotype 12); SEQ ID NO: 187 TCACAAATCACTCTCAAGACAAATCGTGAAAAAGAGCAACCACAAGGTCTAGTCTCTGAGCCAACCACAACTG AGCTAGCTGACACAGATGCAGCACCAATGGCTAATACAGGTCCTGATGCGACTCAAAAAAAGCGCTTCTTTACC 15 GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCATCAAAAAACACAAGATCTATC 20 GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA 25 GCAATTGAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAAC ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTaTTCAGAGTCTGTCGActttaaaGACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCaAAGAGTCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAA 30 AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCC GGATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG \mathbf{T}

> Spy0416A-7 / Schmitz 1/55 (serotype 118); SEQ ID NO: 188 TCACAAGTAACTCCAGAGACAAATCGTGAAAAAGAGCAACCACAAGGTCTAGTCTCTGAGCCAACAACAACTG AGCTAGCTGACACAGATGCAGCACCAATGGCTAATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG 40 TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCATCAAAAAACACAAGATCTATC GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT 45 AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCCTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA GCAATTGAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGATCCATTGGCGACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAAC 50 ATCAGTAGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTGGAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTaTTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCAaAGAGTCAACTGATGCGGGTTATAACGCACAAAACGTTAAAGGTAA AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCC CTGGGAGTACTTATTTTAATAACAAGCCTGGTCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGG 55 GGATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG T

> Spy0416A-7 / Schmitz 1/56 (serotype 28); SEQ ID NO: 189
TCACAAATCACTCCCAAGATAAATCGTGAAAAAGAGCAACCACAAGGTCTAGTCTCTGAGCCAACCACAACTG

AGCTAGCTGACACAGATGCAGCACCAATGGCTAATACAGGTCCTGATGCGACTCAAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGGTACAAAGGACAAGGT AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT 5 GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGCCAAAAAGCCATCAAAAAAACACAAGATCTATCGTCCCC AATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGAC ACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGTAAA GAAGCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGACCCAAGTCATGTTCATGCGTGTTTTTTG CCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCCTTAGGAGCAGA 10 TGTGATCAACCTGAGTCTTGGGACCGCTAATGGTGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATT GAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATG ATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGT GGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAAACCGTGCCGATTTA AACCATGGTAAAGCCATCTaTTCAGAGTCTGTCGACTTtAAAGACATAAAAGATAGCCTAGGTTATGATAAAT 15 CGCATCAATTTGCTTATGTCAAAGAGTCAACTGATGCGGGTTATAACGCACAAGACGTTAAAAGGTAAAATTGC TTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGA GTACTTATTTTTAATAACAAGCCTGGTCAATCAAACCGCTCAATGCGCCTAACAGCTAATGGGATGGGGATAC CATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGT

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> Spy0416A-7 / Schmitz 1/94 (serotype 1); SEQ ID NO: 190 TCACAAATCACTCTCAAGACAAATCGTGAAAAAGAGCAATCACAAGATCTAGTCTCTGAGCCAACCACAACTG AGCTAGCTGACACAGATGCAGCATCAATGGCTAATACAGGTTCTGATGCGACTCAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG 25 TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCCATCAAAAAAACACAAGATCTATC GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT 30 AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACATCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA GCAATTGAAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGATCCATTGGCGACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAAC 35 ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTATTCAGAGTCTGTCGACTttAAAAGACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCaAAGAGTCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAA AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCT 40 GGATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG \mathbf{T}

> Spy0416A-7 / Schmitz 1/253 (serotype 49); SEQ ID NO: 191 TCACAAGTAACTCCAGAGACAAATCGTGAAAAAGAGCAACCACAAGGTCTAGTCTCTGAGCCAACAACAACTG 45 AGCTAGCTGACACAGATGCAGCACCAATGGCTAATACAGGTCCTGATGCGACTCAAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT 50 GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGATGCAGAGCCAAAAAGCCCATCAAAAAAACACAAGATCTATC GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACGTCATGGGATÇAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG 55 AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA GCAATTGAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAAC ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGGATTAGAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATG 60

- > Spy0416A-7 / Schmitz 1/174 (serotype 6); SEQ ID NO: 192 TCACAAATCACTCCCAAGACAAATCGTGAAAAAGAGCAATCACAAGATCTAGTCTCTGAGCCAACCACAACTG AGCTAGCTGACACAGATGCAGCATCAATGGCTAATACAGGTCCTGATGCGACTCAAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC 10 AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCCATCAAAAAAACACAAGATCTATC GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA 15 CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA GCAATTGAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG 20 ACCATGATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAAC ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTaTTCAGAGTCTGTCGACTTTAAAAAACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCAAaGAGTCAACTGATGCGGGTTATAACGCACAAGACGTTAAAGGTAA AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCC 25 GAATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG T
- > Spy0416A-7 / Schmitz 1/176 (serotype 83); SEQ ID NO: 193 30 TCACAAATCACTCTCAAGACAAATCGTGAAAAAGAGCAACCACAAGGTCTAGTCTCTGAGCCAACCACAACTG AGCTAGCTGACACAGATGCAGCACCAATGGCTAATACAGGTCCTGATGCGACTCAAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG 35 TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGAGGATTTT GATGAGGACTGGGAAAACTTTGAGTTTGATGCAGAGGCAGAGCCAAAAAGCCCATCAAAAAAACACAAGATCTATC GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG 40 TTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA GCAATTGAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCCAAC ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTAGAAAACCGTGCC 45 GATTTAAACCATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAAAACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCAAAGAGTCAACTGATGCGGGTTATAAAGCACAAGACGTTAAAGGTAA AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCC CTGGGAGTACTTATTTTAATAACAAGCCTGGTCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGG GGATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG 50 \mathbf{T}

GTCCCCAATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGA CTGGACACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAAT AGCAAAGAAGCCGCTGCTACTGGAGAACGCTTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTG TTTTTGCCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGG AGCAGATGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAA 5 GCAATTGAAAAAAGCTAAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTG ACCATGATGACCCATTGGCAACAAATCCAGACTATGGTTTGGTTTGGTTCTCCCTCAACAGGTCGAACACCAAC ATCAGTGGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGTCAAAGAATTGGAAAACCGTGCC GATTTAAACCATGGTAAAGCCATCTaTTCAGAGTCTGTCGACTTtAAAGACATAAAAGATAGCCTAGGTTATG ATAAATCGCATCAATTTGCTTATGTCAAAGAGTCAACTGATGCGGGTTATAAAGCACAAGACGTTAAAAGATAA 10 AATTGCTTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCC GGATACCATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAG T

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> Spy0416A-7 / Schmitz 1/22 (serotype 4); SEQ ID NO: 195 TCACAAATCACTCCCAAGACAAATCGTGAAAAAGAGCAACCACAAGGTCTAGTCTCTGAGCCAACCACAACTG AGCTAGCTGACACAGATGCAGCATCAATGGCTAATACAGGTCCTGATGCGACTCAAAAAAGCGCTTCTTTACC GCCAGTCAATACAGATGTTCACGATTGGGTAAAAACCAAAGGAGCTTGGGACAAGGGATACAAAGGACAAGGC AAGGTTGTCGCAGTTATTGACACAGGGATCGATCCGGCCCATCAAAGCATGCGCATCAGTGATGTATCAACTG TGATAAAGTTGTTTTTGCACATAATTATGTGGAAAATAGCGATAATATCAAAGAAAATCAATTCGGGGATTTT AATCAACCCAGGCACCGAAAGAAACTGTTATCAAAACAGAAGAAACAGATGGTTCACATGATATTGACTGGAC ACAAACAGACGATGACACCAAATACGAGTCACACGGTATGCATGTGACAGGTATTGTAGCCGGTAATAGCAAA GAAGCCGCTGCTACTGGAGAACGCTTTTTAGGAATTGCACCAGAGGCCCAAGTCATGTTCATGCGTGTTTTTG CCAACGACGTCATGGGATCAGCTGAATCACTCTTTATCAAAGCTATCGAAGATGCCGTGGCTTTAGGAGCAGA TGTGATCAACCTGAGTCTTGGAACCGCTAATGGGGCACAGCTTAGTGGCAGCAAGCCTCTAATGGAAGCAATT GAAAAAGCTAAAAAAGCCGGTGTATCAGTTGTTGTAGCAGCAGGAAATGAGCGCGTCTATGGATCTGACCATG ATGATCCATTGGCAACAAATCCAGACTATGGTTTGGTCGGTTCTCCCTCAACAGGTCGAACACCAACATCAGT GGCAGCTATAAACAGTAAGTGGGTGATTCAACGTCTAATGACGGCCAAAGAATTAGAAAACCGTGCCGATTTA AACCATGGTAAAGCCATCTATTCAGAGTCTGTCGACTTTAAAGACATAAAAGATAGCCTAGGTTATGATAAAT CGCATCAATTTGCTTATGTCAAAGAGTCAACTGATGCGGGTTATAAAGCACAAGACGTTAAAAGATAAAATTGC TTTAATTGAACGTGATCCCAATAAAACCTATGACGAAATGATTGCTTTGGCTAAGAAACATGGAGCCCTGGGA GTACTTATTTTTAATAACAAGCCTGGTCAATCAAACCGCTCAATGCGTCTAACAGCTAATGGGATGGGGATAC CATCTGCTTTCATATCGCACGAATTTGGTAAGGCCATGTCCCAATTAAATGGCAATGGTACAGGAAGT

4. Spy0872

40 4.1 Full length Spy0872

> Spy0872 / SF370 (serotype 1); SEQ ID NO: 196 GATCAAGTTGATGTGCAATTCCTTGGCGTCAATGATTTTCACGGCGCTCTTGATAATACCGGAACAGCTTACA CACCAAGTGGTAAAATACCAAATGCTGGGACGGCTGCTCAATTAGGTGCTTATATGGATGACGCTGAGATAGA CTTCAAGCAAGCAAATCAAGACGGAACAAGTATACGTGTTCAAGCTGGAGATATGGTCGGAGCCAGTCCTGCT 45 AACTCTGCACTTTTACAAGATGAGCCTACTGTCAAAGTCTTTAACAAAATGAAATTTGAATATGGCACTCTTG GTAATCATGAATTTGACGAAGGACTAGATGAATTTAACCGTATCATGACAGGTCAAGCGCCTGATCCTGAATC AACAATTAATGATATCACCAAACAATATGAGCACGAAGCTTCGCATCAAACCATCGTCATTGCTAATGTTATT GATAAAAAAACCAAGGATATCCCCTATGGTTGGAAACCTTATGCTATAAAAGACATAGCCATTAATGACAAAA TCGTTAAGATTGGCTTCATTGGTGTTGTGACTACAGAGATTCCAAATCTCGTTTTAAAGCAAAACTATGAACA 50 CTATCAATTTTTAGATGTAGCTGAAACCATTGCCAAATATGCTAAAGAACTACAAGAACAACATGTTCATGCT ATTGTGGTTTTAGCTCATGTTCCTGCAACAAGTAAAGATGGTGTTGTTGATCATGAAATGGCTACGGTTATGG AAAAAGTGAACCAAATCTATCCCGAACATAGCATTGATATTATTTTTTGCAGGACATAATCATCAATACACTAA TGGAACTATCGGTAAAACACGTATCGTTCAAGCCCTCTCTCAAGGAAAAGCTTATGCAGATGTCCGTGGTACG CTAGATACTGATACCAATGATTTTATTAAAACTCCATCAGCAAATGTTGTTGCTGTAGCACCAGGTATCAAAA 55 CAGAAAATTCAGATATCAAAGCTATAATAAATCATGCTAATGATATTGTTAAAAACAGTTACTGAACGAAAAAT CGGAACTGCAACTAATTCTTCAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTA GCAACAACGGCTCAGCTTACTATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTG GTATTCGAAGTGACCTAGTTGTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATT TGGTAATATCCTTCAAGTCATTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAA 60

AACCAGACCTATTTTCTTCAAATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATA CCCCCTTCAAGATAGTTAAGGTTTATAAAGACAATGGTGAAGAAATTAACTTAACAACTACTTACACCGTTGT TGTCAACGACTTTCTTTATGGTGGTGGTGATGGCTTTTCAGCATTTAAAAAAAGCTAAATTAATCGGAGCTATT AACACAGATACTGAAGCTTTCATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTA TAAAAGGGGTTAAAAATTATGTAACTTCAAACCTTGAAAGTTCGACAAAAGTTAATAGTGCTGGTAAACACAG TATCATTAGTAAGGTTTTTTAGAAATCGTGATGGCAATACAGTGTCTAGTGAAGTCATTTCAGACCTTTTTGACT TCTACTGAAAACACTAATAACAGCCTTGGCAAAAAAGAAACAACAACAACAAAAAAATACTATCTCTAGTTCCA CTCTTCCAATAACA

Antigenic fragment Spy0872-2 10 4.2

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> Spy0872-2 / SF370 (serotype 1); SEQ ID NO: 17 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGCAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCCTTCAAGATAGTTAAG GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAGCTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT 20 CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAAGGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGTTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTA GAAATCGTGATGGCAATACAGTGTCTAGTGAAGTCATTTCAGACCTTTTTGACTTCTACTGAAAACACTAATAA CAGCCTTGGCAAAAAAGAAACAACAACAAACAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA

Homologous sequences of other S. pyogenes isolates and/or serotypes 4.3

> Spy0872-2 / Schmitz 1/7 (serotype 4); SEQ ID NO: 197 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACGGCTCAGCTTAC 30 TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGAAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCCTTCAAGATAGTTAAG 35 GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAGCTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAAGGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGTTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTATCATTAGTAAGG TTTTTAGAAATCGTGATGGCAATATAGTGTCTAGTGAAATCATTTCAGACCTTTTGACTTCTACTGAAAACAC TAATAACAGCTTTGGCAAAAAAGAGATAACAACAAACaAAAATACTATCTCTAATTCCACTCTTCCAATAACA 40

> Spy0872-2 / Schmitz 1/39 (serotype 12); SEQ ID NO: 198 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCTTCAAGATAGTTAAG GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAGCTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAAGGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGCTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTTA GAAATCGTGATGGCAATATAGTGTCTAGTGAAATCATTTCAGACCTTTTTGACTTCTACTGAAAACACTAATAA CAGCCTTGGCAAAAAAGAAACAACGACAAACAAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA

> Spy0872-2 / Schmitz 1/55 (serotype 118); SEQ ID NO: 199 GCTATAATAAaTCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA

- > Spy0872-2 / Schmitz 1/56 (serotype 28); SEQ ID NO: 200 10 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACAGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGATCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA 15 AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCCTTCAAGATAGTTAAG CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGTTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTA 20 GAAATCGTGATGGCAATATAGTGTCTAGTGAGATCATTTCAGACCTTTTTGACTTCTACTGAAAACACTAATAA CAGCCTTGGCAAAAAAGAAACAACAACAAACAAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA
- > Spy0872-2 / Schmitz 1/94 (serotype 1); SEQ ID NO: 201 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT 25 CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGCAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCCTTCAAGATAGTTAAG 30 GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAGCTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAGGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGTTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTA GAAATCGTGATGGCAATACAGTGTCTAGTGAAGTCATTTCAGACCTTTTTGACTTCTACTGAAAACACTAATAA 35 CAGCCTTGGCAAAAAAGAAACAACAACAAACAAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA
- > Spy0872-2 / Schmitz 1/253 (serotype 49); SEQ ID NO: 202 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACAGCTCAGCTTAC 40 TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGATCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATTCACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCCTTCAAGATAGTTAAG 45 CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAGGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGCTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTTA GAAATCGTGATGGCAATATAGTGTCTAGTGAAATAATTTCAGACCTTTTTGACTTCTACTGAAAAACACTAATAA CAGCCTTGGCAAAAAAAAAACAACGACaAACAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA 50

5 > Spy0872-2 / Schmitz 1/177 (serotype 22); SEQ ID NO: 204 GCTATAATAAATCATGCTAATGATATTGTTAAAAACAGTTACTGAACGAAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA 10 TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCTTCAAGATAGTTAAG GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAAGCTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAGGGGGTTAAAAATTAT 15 GTAACTTCAAACCTTGAAAGCTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTA GAAATCGTGATGGCAATATAGTGTCTAGTGAAATCATTTCAGACCTTTTTGACTTCTACTGAAAAACACTAATAA CAGCCTTGGCAAAAAAGAAACAACAAACAAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA

- > Spy0872-2 / Schmitz 1/234 (serotype 44); SEQ ID NO: 205 20 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGGAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA 25 AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCTTCAAGATAGTTAAG GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAAACTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAAGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGCTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTAGTAAGGTTTTTA 30 GAAATCGTGATGGCAATATAGTGTCTAGTGAAATCATTTCAGACCTTTTTGACTTCTACTGAAAACACTAATAA CAGCCTTGGCAAAAAAGAAACAACGACaAACAAAAATACTATCTCTAGTTCCACTCTTCCAATAACA
- > Spy0872-2 / Schmitz 1/22 (serotype 4); SEQ ID NO: 206 GCTATAATAAATCATGCTAATGATATTGTTAAAACAGTTACTGAACGAAAAATCGGAACTGCAACTAATTCTT 35 CAACTATTTCTAAAACAGAAAATATTGATAAAGAATCTCCTGTCGGTAACTTAGTAACAACGGCTCAGCTTAC TATTGCTAAGAAAACTTTTCCAACTGTTGACTTTGCTATGACCAATAATGGTGGTATTCGAAGTGACCTAGTT GTCAAAAATGACCGGACCATCACCTGGGAAGCTGCACAGGCTGTACAACCATTTGGTAATATCCTTCAAGTCA TTCAAATGACTGGTCAACACATTTACGATGTCCTAAATCAGCAATACGATGAAAACCAGACCTATTTTCTTCA AATGTCAGGTTTAACATACACTTATACAGATAATGATCCTAAGAACTCTGATACCCCCTTCAAGATAGTTAAG 40 GTGGTGGTGATGGCTTTTCAGCATTTAAAAAAAGCTAAATTAATCGGAGCTATTAACACAGATACTGAAGCTTT CATCACATATATCACAAATTTAGAAGCATCAGGTAAAACTGTTAATGCTACTATAAAAAGGGGGTTAAAAATTAT GTAACTTCAAACCTTGAAAGTTCGACAAAAGTTAATAGTGCTGGTAAACACAGTATCATTATCATTAGTAAGG TTTTTAGAAATCGTGATGGCAATATAGTGTCTAGTGAAATCATTTCAGACCTTTTGACTTCTACTGAAAACAC 45

5. Further Sequences

CTGGTAAATTCAAACGAGACCAGAGCCTTAACGAGCATGAAGCTATGGTAAATGTCGAACCACTGTCTCATGT AGTAAAAGACAATATAAAGGTCATAGATGAAAAATCAACAGGGCGGTTTGAGCCTTTTAGACCTAATGAAGAT GAGAAGGAGAAGCCTGCCAGCGATGTTAAGGTAAGACCAGCAGAAGTTGGTAGCTGGCTAGAACCAGCGACAG CTCTTCCTAGTGTTGAAATGAGCGCTGAGGACAGGTTAAAAAAGT

> Spy1536 / SF370 (serotype 1); SEQ ID NO: 207 ATTGAAATGCCTGGAGGCGCTTACGATATTCGGACTGTCTTACAAGTCAATGGCAAAGAAGACAAACGAAAAG 20 GAGCTTACCAGTTTGTTGCAGTGGGCATTAGTCGTGCCAGCCTCGCTCAGCTATTATATGCTTGGCTGACACC GTTTACTGAAATTAGTACAGCAGAAGATACAACAGGCGGATACAGCGATGCTGATTTCCTTCGAATTAATCAA TTTTACATGGAAACATCACAAAATGCAGCTATTTATCAAGCTTTATCCTTAGCTGGAAAACCAGTTACATTAG ATTATAAAGGCGTATATGTTTTAGACGTAAACAACGAATCTACTTTTAAAGGAACGCTACACTTAGCAGATAC TGTAACAGGTGTAAATGGTAAACAGTTTACTAGTTCAGCAGAACTTATTGACTATGTTTCTCACCTAAAACTA 25 GGGGATGAAGTTACGGTTCAGTTTACGAGTGATAATAAGCCTAAAAAAAGGAGTTGGCCGTATTATCAAACTGA AAAATGGGAAAAATGGGATTGGCATTGCCTTGACTGATCATACAAGTGTCAATTCAGAAGACACAGTGATCTT GAAGATTTACGCAAGGGCCGTACAATTGCAGGTACAGGAACTATTGGCAAGGATGGCGAAGTAGGAGATATTG 30 TGATAAGGAAATTAAAAAAGTTAATCCAAATGCTATAAGTAATTACGAAGAAGCCAAACGGGCAGCCAAACGA CTAAAGACCAAAATGAAGATTGTTCCTGTTACGACTGTTCAAGAGGCACTGGTTTATCTTCGCAAA

> Spy1666 / SF370 (serotype 1); SEQ ID NO: 208 35 ACAAAAGAATTTCATCACGTGACCGTACTCCTTCACGAAACAGTGGACATGCTTGACATAAAGCCTGATGGGA TTTATGTTGATGCGACGCTAGGTGGCTCAGGCCACTCAGCTTATTTGTTGTCCAAACTTGGTGAAGAAGGGCA CCTCTATTGTTTTGACCAAGACCAAAAGGCTATTGACAATGCACAAGTTACCCTCAAATCTTATATTGACAAA GGACAGGTAACTTTTATTAAAGATAATTTTAGACACCTCAAAGCACGTTTAACAGCGCTTGGAGTTGATGAAA TTGATGGTATCTTATATGACCTTGGTGTTTCCAGCCCGCAATTGGATGAAAGAGAACGAGGGTTTTCTTATAA ACAAGATGCTCCATTGGATATGCGCATGGATCGTCAGTCGCTCTTAACAGCTTACGAAGTGGTGAATACCTAT 40 CCATTCAATGATTTGGTTAAGATTTTTTTCAAATATGGTGAAGATAAATTCTCCAAGCAGATCGCTCGAAAAA TTGAACAAGCAAGAGCTATTAAGCCTATTGAGACAACAACAGAGTTGGCAGAATTGATTAAGGCAGCAAAGCC AGCTAAAGAGTTGAAGAAAAAAGGCCACCCTGCTAAACAGATTTTTCAAGCTATTCGCATTGAAGTCAATGAT GAATTGGGAGCGCCGATGAATCTATTCAGGACGCTATGGAATTATTAGCCCTTGATGGTCGTATCTCAGTTA TTACCTTCCATTCTCTGGAAGATCGCCTAACCAAGCAGTTGTTTAAAGAAGCTAGTACGGTGGATGTGCCAAA 45 AGGGCTTCCTCTAATTCCTGAAGATATGAAACCTAAGTTTGAACTTGTTTCACGTAAGCCGATCTTACCTAGT CATTCAGAGTTAACAGCTAATAAAAGGGCACACTCAGCCAAGCTACGTGTTGCCAAAAAAATTCGGAAA

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Claims

- A peptide consisting of one antigen of *S. pyogenes* of the SEQ ID NO: 4, SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 7, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 3 or a functional active variant of one antigen of *S. pyogenes* of the SEQ ID NO: 4, SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 7, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 3.
- 2. A peptide consisting of one antigen of *S. pyogenes* of the SEQ ID NO: 4, SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 7, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 3 or a functional active variant of one antigen of *S. pyogenes* of the SEQ ID NO: 4, SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 7, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 3, and
 - a) 1 to 350 additional amino acid residue(s), preferably 1 to 200, more preferably 1 to 150, even more preferably at most 1 to 100, still more preferably at most 1 to 50, most preferably 1, 2, 3, 4, 5, 10, 15, 20 or 25 additional amino acids residue(s) if the antigen is SEQ ID NO: 1; or
 - b) 1 to 200 additional amino acid residue(s), preferably 1 to 150, more preferably 1 to 100, even more preferably at most 1 to 50, still more preferably at most 1 to 25, most preferably 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 additional amino acids residue(s) if the antigen is SEQ ID NO: 2; or
 - c) 1 to 100 additional amino acid residue(s), preferably 1 to 75, more preferably 1 to 50, even more preferably at most 1 to 25, still more preferably at most 1 to 10, most preferably 1, 2, 3, 4 or 5 additional amino acids residue(s) if the antigen is that of SEQ ID NO: 3; or
 - d) 1 to 150 additional amino acid residue(s), preferably 1 to 100, more preferably 1 to 75, even more preferably at most 1 to 50, still more preferably at most 1 to 25, most preferably 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 additional amino acids residue(s) if the antigen is that of SEQ ID NO: 4; or
 - e) 1 to 450 additional amino acid residue(s), preferably 1 to 300, more preferably 1 to 150, even more preferably at most 1 to 100, still more

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- f) 1 to 250 additional amino acid residue(s), preferably 1 to 200, more preferably 1 to 150, even more preferably at most 1 to 100, still more preferably at most 1 to 50, most preferably 1, 2, 3, 4, 5, 10, 15, 20 or 25 additional amino acids residue(s) if the antigen is SEQ ID NO: 6 or SEQ ID
- 3. The peptide of any of claims 1 or 2 further consisting of at least one amino acid residue heterologous to the antigen, preferably an additional amino acid sequence comprising a marker protein.

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- 4. The peptide of any of claims 2 or 3, wherein the additional amino acid residue(s) is/are flanking the antigen C-terminally, N-terminally or C- and N-terminally.
- 5. The peptide of any of claims 1 to 4, wherein the functional active variant is essentially identical to any of the antigens of the SEQ ID NO: 4, SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 7, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 3, but differs from the antigens of any of the SEQ ID NO: 4, SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 7, SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 3 in that it is derived from a homologous sequence of a different serotype of *S. pyogenes*, particularly wherein the serotype is M2, M3, M4, M5, M6, M11, M12, M14, M19, M22, M24, M25, M28, M44, M49, M57, M59, M60, M61, M76, M83, M84, M87, M89 or M118.
- 6. The peptide of any of claims 1 to 5, wherein the functional active variant is a portion of any of the SEQ ID NOS: 1 to 7 consisting of at least 60%, preferably at least 70%, more preferably at least 80%, still more preferably at least 90%, even more preferably at least 95%, most preferably 99% of the amino acids of the antigen of any of the SEQ ID NOS: 1 to 7.
- 7. The peptide of any of claims 1 to 6, wherein the functional active variant of the antigen of any of the SEQ ID NOS: 1 to 7 has at least 50% sequence identity to the

antigen of any of the SEQ ID NOS: 1 to 7, especially at least 60%, preferably at least 70%, more preferably at least 80%, still more preferably at least 90%, even more preferably at least 95%, most preferably 99% sequence identity to the antigen of any of the SEQ ID NOS: 1 to 7.

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- 8. The peptide of claim 7, wherein the variant is derived from the antigen of any of the SEQ ID NOS: 1 to 7 by at least one conservative amino acid substitution.
- 9. A peptide comprising an amino acid sequence with at least 95% sequence identity to at least one of SEQ ID NO: 1, 2, 3, 4, 5, 6 or 7, wherein said peptide is not Spy0269, Spy0292, Spy0416A, or Spy0872.
 - 10. A peptide characterized in that it comprises at least 2, preferably at least 3, more preferably at least 4 antigens as defined in any of claims 1 to 9.

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- 11. A nucleic acid coding for the peptide according to any of claims 1 to 10 or a nucleic acid complementary thereto, particularly a DNA sequence of any of the sequences of SEQ ID NOS: 11 to 17 or the corresponding RNA sequence.
- 20 12. The nucleic acid of claim 11, wherein the nucleic acid is located in a vector.
 - 13. A pharmaceutical composition, especially a vaccine, comprising
 - (i) at least one peptide according to any of claims 1 to 10 and/or
 - (ii) at least one peptide comprising or consisting of the sequence of any of the SEQ ID NO: 8, SEQ ID NO: 9, or SEQ ID NO: 10, or a functional active variant thereof, and
 - (iii) optionally a pharmaceutically acceptable carrier or excipient.
 - 14. A pharmaceutical composition containing
- (i) a nucleic acid according to claim 11 and/or a nucleic acid complementary thereto and/or
 - (ii) a nucleic acid coding for the peptide comprising or consisting of the sequence of any of the SEQ ID NO: 8, SEQ ID NO: 9, or SEQ ID NO: 10,

particularly a DNA sequence of any of the SEQ ID NO: 18, SEQ ID NO: 19, or SEQ ID NO: 20, or a functional active variant thereof or a nucleic

acid complementary thereto or the corresponding RNA sequence, and

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(iii) optionally a pharmaceutically acceptable carrier or excipient.

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- 15. The pharmaceutical composition of claim 14, wherein the nucleic acid is comprised in a vector and/or a cell.
- 16. An antibody or functional active fragment thereof which binds specifically to the antigen of claim 1.
 - 17. The antibody or functional active fragment thereof of claim 16, wherein the antibody is a monoclonal, polyclonal, chimeric or humanized antibody, or wherein the functional active fragment comprises a Fab fragment.
 - 18. A hybridoma cell line which produces the antibody according to claim 16 or 17.
 - 19. A method for producing an antibody according to claim 16 or 17, characterized by the following steps:
 - (a) administering an effective amount of the peptide according to any of claims 1 to 10 to an animal; and
 - (b) isolating the antibody produced by the animal in response to the administration of step (a) from the animal.
- 25 20. A method for producing an antibody according to claim 16 or 17, characterized by the following steps:
 - (a) contacting a B cell with an effective amount of the peptide according to any of claims 1 to 10;
 - (b) fusing the B cell of step (a) with a myeloma cell to obtain a hybridoma cell; and
 - (c) isolating the antibody produced by the cultivated hybridoma cell.

- 21. The method of claim 19 or 20, wherein the isolated antibody is additionally purified.
- 22. A pharmaceutical composition, especially a vaccine, comprising the antibody according to claim 16 or 17.
 - 23. A pharmaceutical composition comprising the peptide as defined in claim 13 or the nucleic acid as defined in claim 14 or the antibody or functional fragment thereof according to claim 16 or 17 for the immunization of a subject against an infection or the treatment of a subject having an infection, wherein the infection is preferably a *S. pyogenes* infection.

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- Use of the peptide as defined in claim 13 or the nucleic acid as defined in claim 14 or the antibody or functional fragment thereof according to claim 16 or 17 for the manufacture of a medicament for immunization against or treatment of an infection, preferably a *S. pyogenes* infection.
- 25. Method of immunizing a subject against an infection or treating a subject having an infection, the method comprising
- 20 (a) administering to the patient an effective amount of the peptide as defined in claim 13 or the nucleic acid as defined in claim 14 or the antibody or functional fragment thereof according to claim 16 or 17.
 - 26. The method of claim 25, wherein the infection is a S. pyogenes infection.
 - 27. A method of diagnosing a S. pyogenes infection comprising the steps of:
 - (a) contacting a sample obtained from a subject with the peptide according to any of claims 1 to 10; and
 - (b) detecting the presence of an antibody against S. pyogenes in the sample.
 - 28. A method of diagnosing a S. pyogenes infection comprising the steps of:
 - (a) contacting a sample obtained from a subject with the antibody according to claim 16 or 17; and

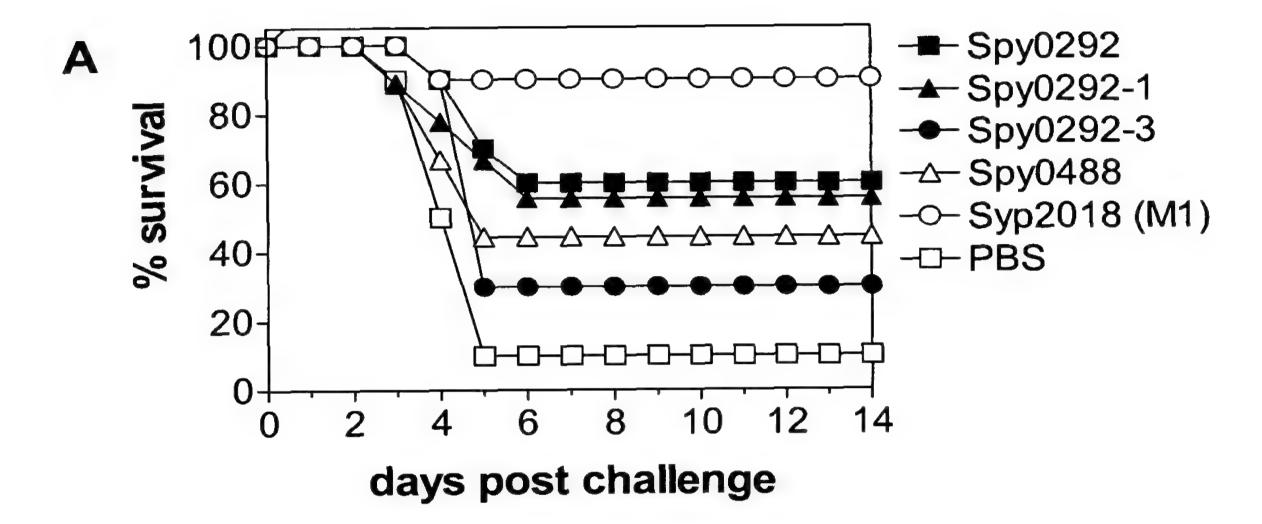
- (b) detecting the presence of an antigen of S. pyogenes in the sample.
- 29. A method for identifying a ligand capable of binding to a peptide according to any of claims 1 to 10 comprising:
 - (a) providing a test system comprising the peptide,

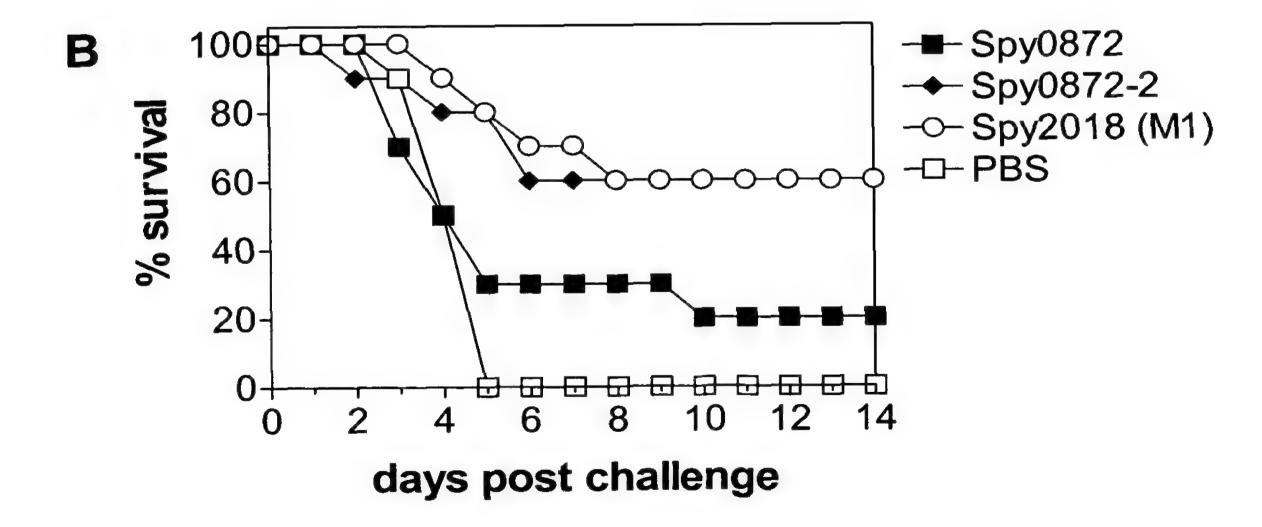
- (b) contacting the test system with a test compound, and
- (c) detecting a signal generated in response to the binding of the test compound to the peptide or functional active variant.
- 10 30. Use of any of the peptide according to any of claims 1 to 10 for the isolation and/or purification and/or identification of an interaction partner of the peptide.

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Figure 1

CFA/IFA model

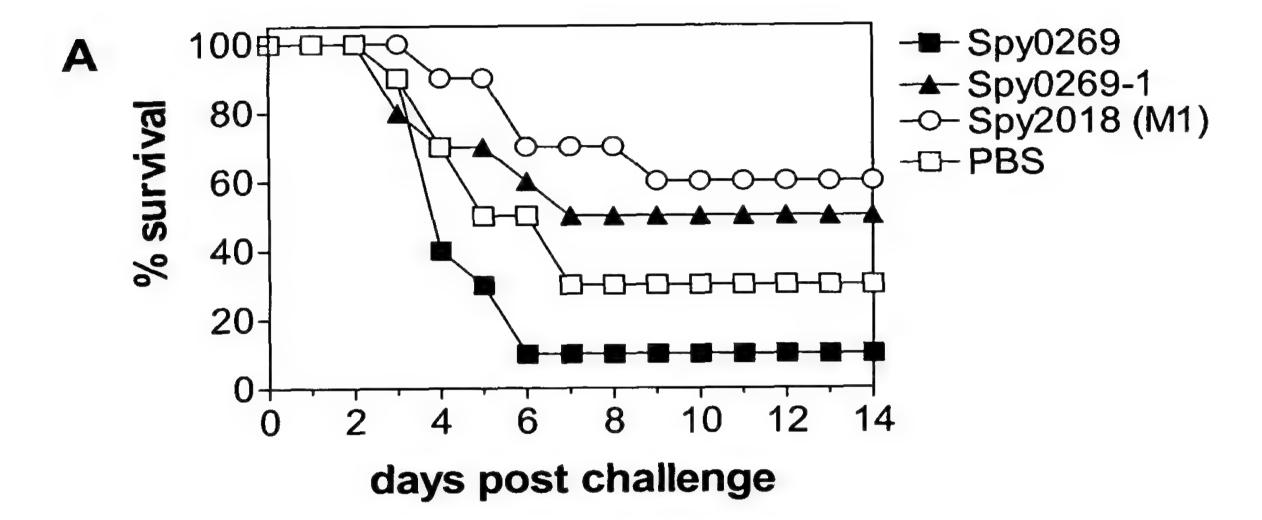


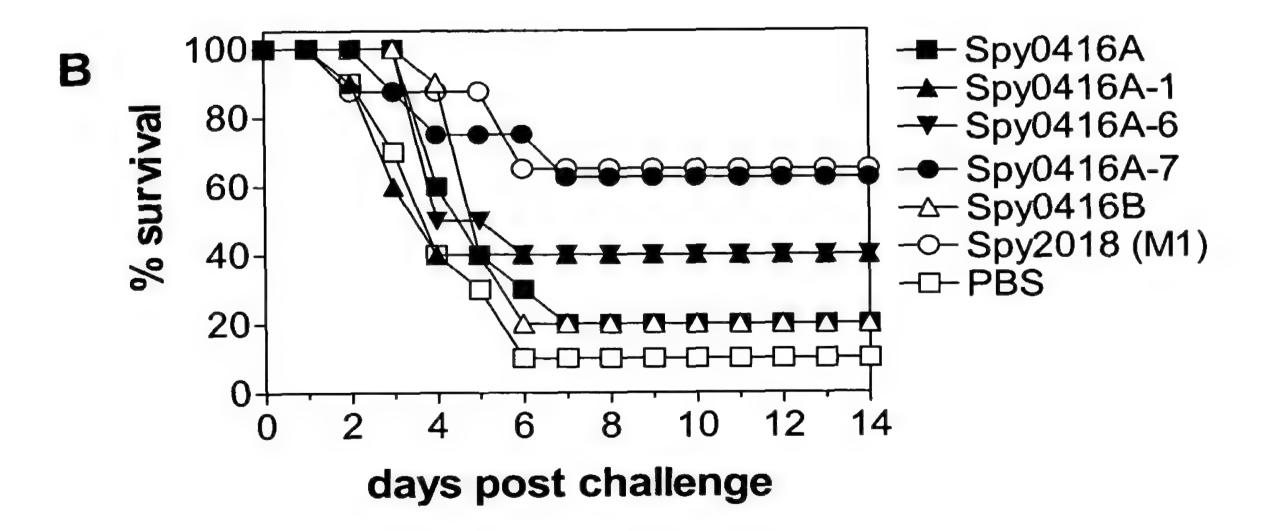


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Figure 2

CFA/IFA model

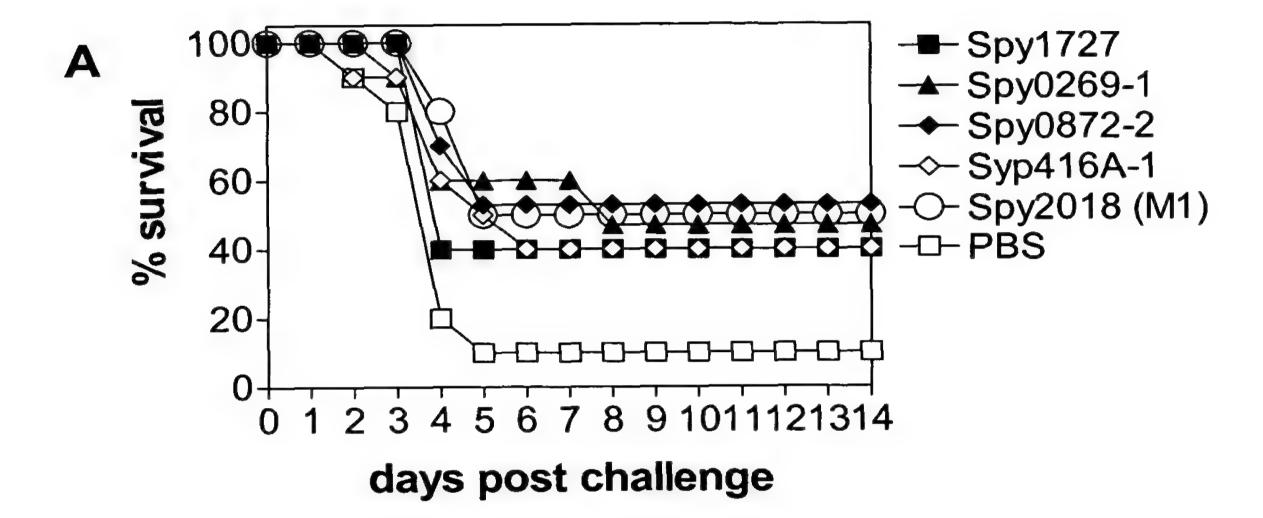


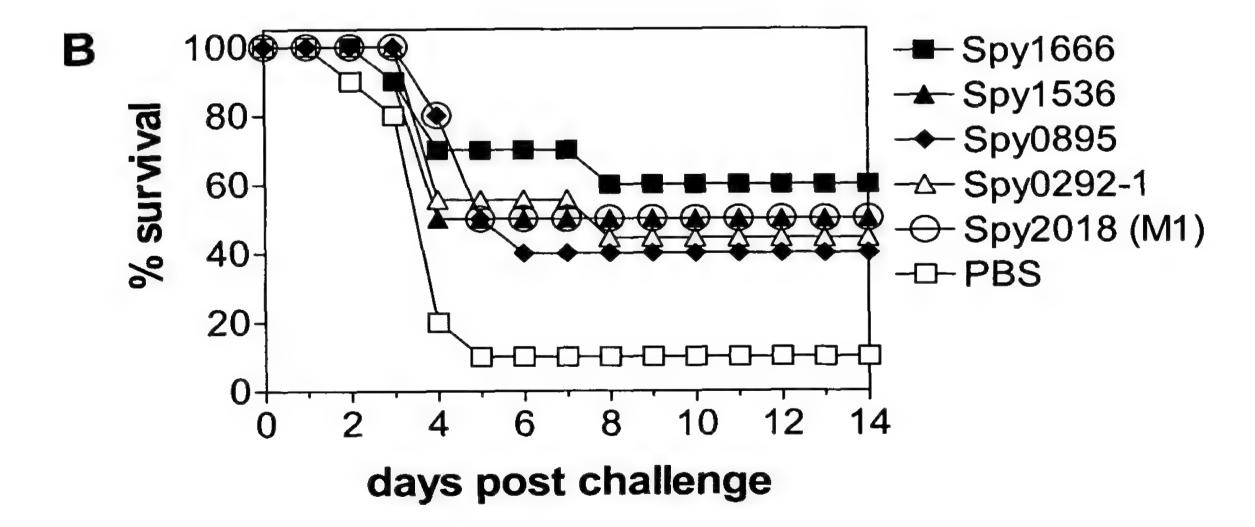


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Figure 3

ALUM model

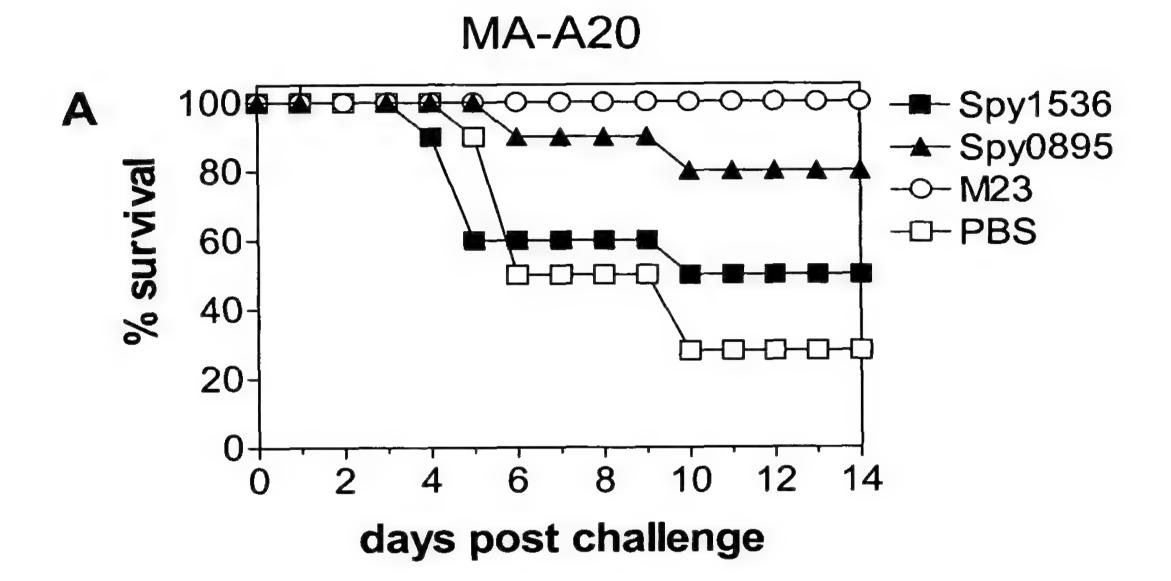


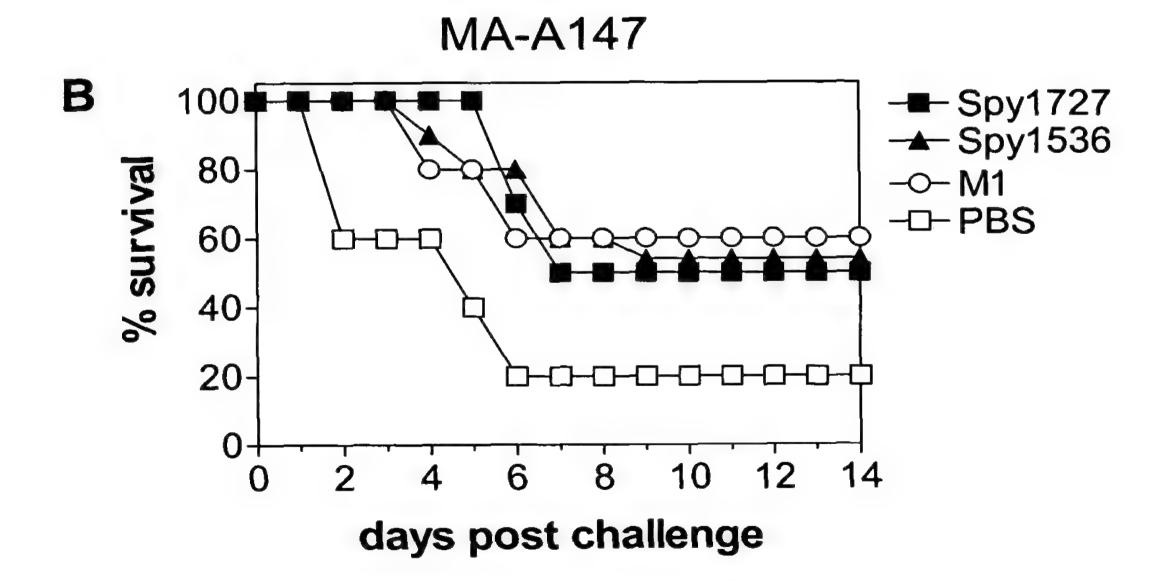


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Figure 4

IC31 i.n. model





INTERNATIONAL SEARCH REPORT

International application No PCT/EP2007/006027

A. CLASSIFICA INV. CO7	TION OF SUBJECT MATTER 7K 16/00	:	·			
According to Inte	ernational Patent Classification (IPC) or to both national class	ification and IPC	,			
B. FIELDS SEARCHED						
Minimum docum CO7K	entation searched (classification system followed by classific	cation symbols)	,			
Documentation s	searched other than minimum documentation to the extent the	at such documents are included in the fields se	arched			
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)						
EPO-Inter	rnal, WPI Data					
C. DOCUMENTS	S CONSIDERED TO BE RELEVANT					
	ation of document, with indication, where appropriate, of the	relevant passages	Relevant to claim No.			
			· · · · · · · · · · · · · · · · · · ·			
X	WO 2005/032582 A2 (CHIRON CORP GRANDI GUIDO [US]; TELFORD JOHN BENSI GIULIANO) 14 April 2005 ([US]; 2005-04-14)	1-30			
	page 26, line 23 - page 53, line 25; claims 2,11,27; sequence 122					
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Further d	locuments are listed in the continuation of Box C.	X See patent family annex.	,			
* Special categ	ories of cited documents :	"T" later document published after the inte- or priority date and not in conflict with	rnational filing date			
"A" document d considered	efining the general state of the art which is not it to be of particular relevance	cited to understand the principle or the invention	eory underlying the			
"E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to						
"L" document which may throw doubts on priority claim(s) or involve an inventive step when the document is taken alone which is cited to establish the publication date of another "Y" document of particular relevance; the claimed invention						
citation or o	other special reason (as specified) eferring to an oral disclosure, use, exhibition or	cannot be considered to involve an involve document is combined with one or mo	rentive step when the re other such docu-			
other mear		ments, such combination being obviouin the art.				
later than t	he priority date claimed	*&* document member of the same patent t	amily			
	a) completion of the international search	Date of mailing of the international sear	rch report			
18 3	September 2007	27/11/2007	_			
Name and mailing address of the ISA/ Authorized officer Furn poor Retart Office, B.E. 5818 Retartion 2						
•	European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel (+31-70) 340-2040. Tv. 31 651 eno nl.		•			
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016 Stoyanov, Borislav						

International application No. PCT/EP2007/006027

INTERNATIONAL SEARCH REPORT

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)					
This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:					
1. X Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:					
see FURTHER INFORMATION sheet PCT/ISA/210					
2. Claims Nos.:					
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:					
3. Claims Nos.:					
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).					
Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)					
This International Searching Authority found multiple inventions in this international application, as follows:					
This international Searching Additionly found matuple inventions in this international application, as follows.					
see additional sheet					
1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.					
2. As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.					
3. As only some of the required additional search fees were timely paid by the applicant, this international search reportcovers					
only those claims for which fees were paid, specifically claims Nos.:					
4. X No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:					
1-30 (only partially)					
Remark on Protest The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.					
The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.					
No protest accompanied the payment of additional search fees.					

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box II.1

Although claims 25-26 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

Although claims 27-28 are directed to a diagnostic method practised on the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

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FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 4, or a variant thereof, the nucleic acid encoding it and the uses thereof.

2. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 1, or a variant thereof, the nucleic acid encoding it and the uses thereof.

3. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 2, or a variant thereof, the nucleic acid encoding it and the uses thereof.

4. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 7, or a variant thereof, the nucleic acid encoding it and the uses thereof.

5. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 5, or a variant thereof, the nucleic acid encoding it and the uses thereof.

6. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 6, or a variant thereof, the nucleic acid encoding it and the uses thereof.

7. claims: 1-30 (only partially)

A peptide consisting of SEQ ID NO. 3, or a variant thereof, the nucleic acid encoding it and the uses thereof.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/EP2007/006027

Patent document	Publication	Patent family	Publication
cited in search report	date	member(s)	date
WO 2005032582 A2	14-04-2005	CA 2532369 A1 EP 1648500 A2 JP 2007500726 T	